



US006884903B2

(12) **United States Patent**
Lorenz et al.

(10) **Patent No.: US 6,884,903 B2**
(45) **Date of Patent: Apr. 26, 2005**

(54) **SUBSTITUTED
SULFONYLAMINOMETHYLBENZOIC ACID
(DERIVATIVES) AND THEIR PREPARATION**

(75) Inventors: **Klaus Lorenz**, Weiterstadt (DE);
Hans-Joachim Ressel, Hattersheim
(DE); **Lothar Willms**, Hofheim (DE)

(73) Assignee: **Aventis CropScience GmbH**, Frankfurt
(DE)

(*) Notice: Subject to any disclaimer, the term of this
patent is extended or adjusted under 35
U.S.C. 154(b) by 0 days.

(21) Appl. No.: **10/387,649**

(22) Filed: **Mar. 13, 2003**

(65) **Prior Publication Data**

US 2003/0208087 A1 Nov. 6, 2003

Related U.S. Application Data

(62) Division of application No. 09/910,105, filed on Jul. 20,
2001, now Pat. No. 6,538,150.

(30) **Foreign Application Priority Data**

Jul. 24, 2000 (DE) 100 36 184

(51) **Int. Cl.⁷** **C07C 303/00**

(52) **U.S. Cl.** **560/12; 560/14**

(58) **Field of Search** 560/12, 14

(56) **References Cited**

U.S. PATENT DOCUMENTS

4,372,778 A 2/1983 Levitt
5,463,081 A 10/1995 Ort et al.
5,648,315 A 7/1997 Lorenz et al.
5,925,596 A 7/1999 Lorenz et al.
6,221,809 B1 4/2001 Hacker et al.
6,492,301 B1 12/2002 Hacker et al.

FOREIGN PATENT DOCUMENTS

WO WO 00/44227 8/2000

OTHER PUBLICATIONS

Saari et al, J. Heterocyclic Chemistry, (1986), 23 pp.
1253–1255.*

5–Agrochemicals; pp. 303–304, “Effects of Humactants on
the Pesticide Uptake Through Plant Leaf Surfaces”.

Chemische Berichte, 1957; 90. Jahrg; No. 6, pp. 841–1178.

J. Med. Chem. 1986 29, 585–589, Gelb et al, “Substituted
Isatoic Anhydrides: Selective Inactivators of Trypsin–like
Serine Proteases”.

Dankwardt et al, Bijoorg Med. Lett. 1997, vol. 7, No. 14, pp.
1921–1926.

Saari et al, “A Convenient Synthesis of Nitro–Substituted
1,2–Benzisothiazol–3(2H)–one 1,1 Dioxides (Nitrosacchar-
ins)”, J Heterocyclic Chem, 23, 1986, pp. 1253–1255, also
referred to as XP–002178946.

* cited by examiner

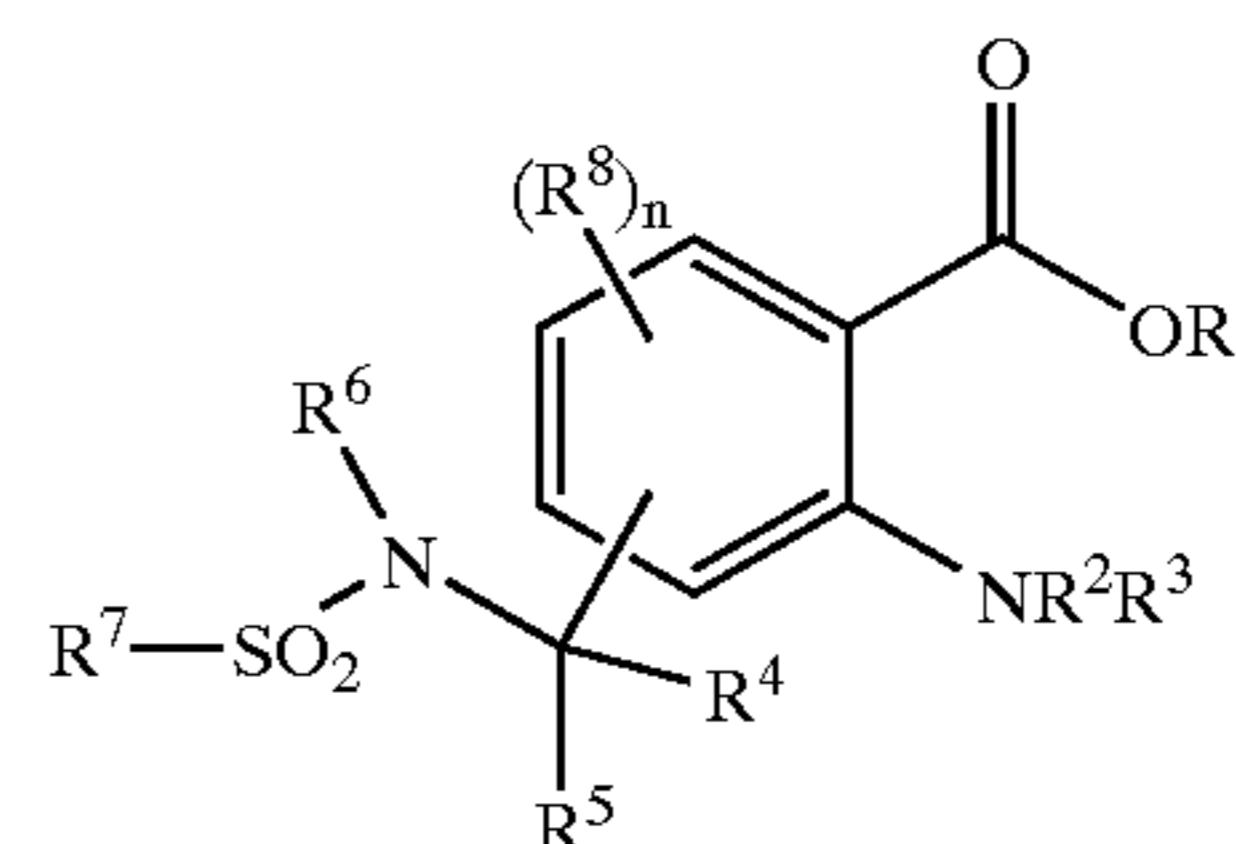
Primary Examiner—Rita Desai

Assistant Examiner—Hector M. Reyes

(74) *Attorney, Agent, or Firm*—Frommer Lawrence &
Haug LLP

(57) **ABSTRACT**

The present invention relates to a compound of the formula
(I)



(I)

The compound according to the invention is suitable for the
preparation of herbicidal active substances such as sulfonyl-
ureas.

9 Claims, No Drawings

1

**SUBSTITUTED
SULFONYLAMINOMETHYLBENZOIC ACID
(DERIVATIVES) AND THEIR PREPARATION**

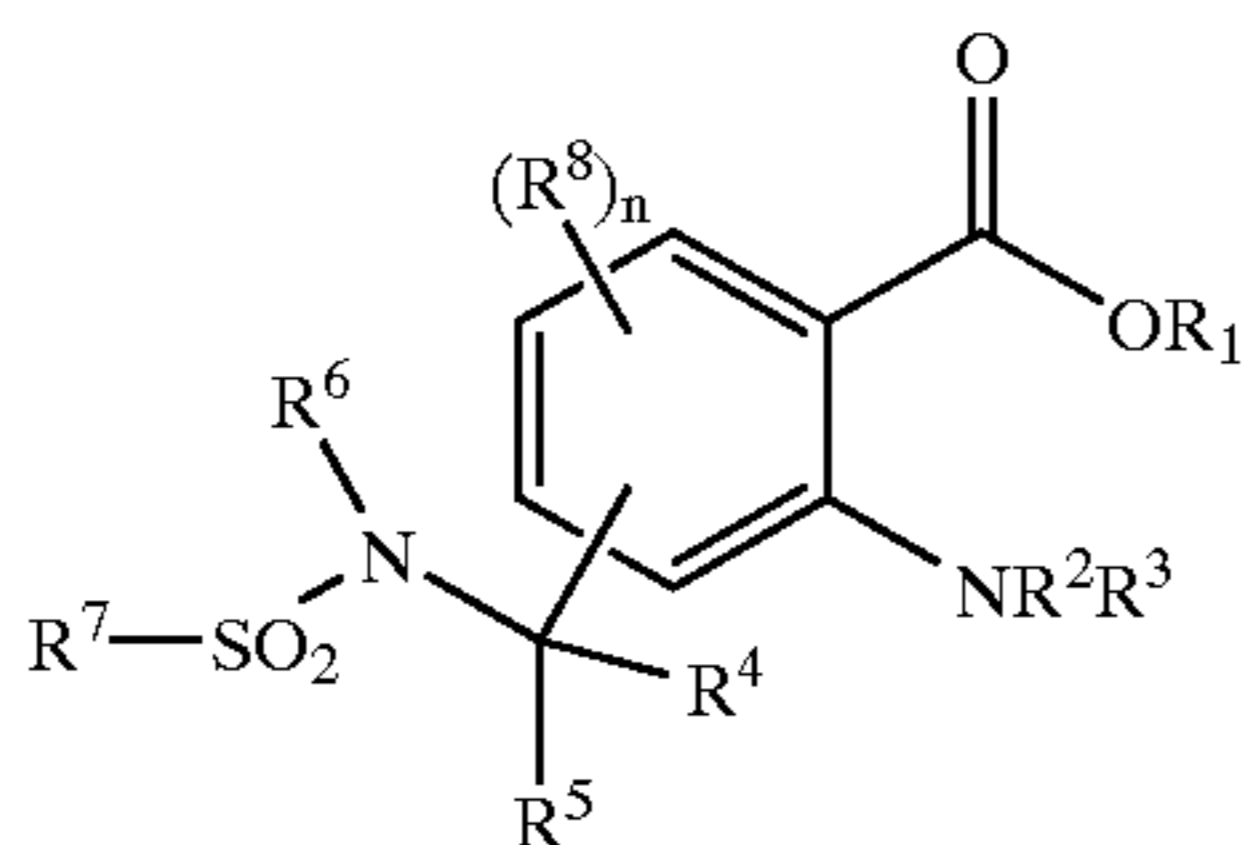
This is a division of application Ser. No. 09/910,105, filed Jul. 20, 2001, now U.S. Pat. No. 6,538,150.

The invention relates to the technical field of the intermediates for the preparation of active substances, in particular herbicidally active sulfonylureas.

It is known that aromatic amines can be reacted to give sulfonic acid derivatives such as sulfochlorides and further to give sulfonamides which, in turn, can be employed for the preparation of herbicidally active sulfonylureas (Meerwein et al., Chem. Berichte 90, 841-852 (1957) and EP-A-574418).

A substituted anthranilic acid is known from J. Med. Chem. 1986, Vol. 29, No. 4, page 585 as intermediate for the preparation of certain anhydrides which are suitable for inactivating trypsin-like enzymes.

It was an object to provide novel chemical compounds which are suitable for the preparation of herbicidally active sulfonylureas. Surprisingly, this object is achieved by compounds of the formula (I)



in which

R^1 is H, (C_1-C_8) alkyl, (C_3-C_8) alkenyl or (C_3-C_8) alkynyl, where the last 3 radicals are unsubstituted or substituted, for example by one or more radicals selected from the group consisting of halogen, (C_1-C_4) alkoxy, (C_1-C_4) alkylthio, $[(C_1-C_4)alkyl]carbonyl$ or $[(C_1-C_4)alkoxy]carbonyl$,

R^2, R^3 independently of one another are H or acyl, preferably H,

R^4, R^5 are H,

R^6 is H or (C_1-C_8) alkyl which is unsubstituted or substituted, for example by one or more radicals selected from the group consisting of halogen, (C_1-C_4) alkoxy, (C_1-C_4) alkylthio, (C_1-C_4) alkylsulfinyl, (C_1-C_4) alkylsulfonyl, $[(C_1-C_4)alkyl]carbonyl$ or CN, preferably H,

R^7 is (C_1-C_8) alkyl, (C_3-C_8) alkenyl or (C_3-C_8) alkynyl which are unsubstituted or substituted, for example by one or more radicals selected from the group consisting of halogen, (C_1-C_4) alkoxy or (C_1-C_4) alkylthio, or R^7 is (C_6-C_{14}) aryl (for example phenyl) which is unsubstituted or substituted, for example by one or more radicals selected from the group consisting of halogen, NO_2 , CN, (C_1-C_4) alkyl, (C_1-C_4) haloalkyl or (C_1-C_4) alkoxy, or R^7 is mono- or di- (C_1-C_8) alkylamino which is unsubstituted or substituted, for example by one or more radicals selected from the group consisting of halogen, (C_1-C_4) alkoxy, (C_1-C_4) alkylthio, (C_1-C_4) alkylsulfinyl, (C_1-C_4) alkylsulfonyl, $[(C_1-C_4)alkyl]carbonyl$, $[(C_1-C_4)alkoxy]carbonyl$ or CN, or

R^6 and R^7 together form a chain of the formula $-(CH_2)_m B_{m1}-$ which is unsubstituted or substituted, for example

2

by one or more (C_1-C_4) alkyl radicals, and where $m=2, 3$ or 4, $m^1=0$ or 1 and $B=CO$ or SO_2 ,

R^8 radicals, are identical or different and are (C_1-C_4) alkyl, (C_1-C_4) alkoxy, $[(C_1-C_4)alkyl]carbonyl$ or $[(C_1-C_4)alkoxy]carbonyl$ which are unsubstituted or substituted, for example by one or more radicals selected from the group consisting of halogen, (C_1-C_4) alkoxy, (C_1-C_4) alkylthio, $[(C_1-C_4)alkyl]carbonyl$ or $[(C_1-C_4)alkoxy]carbonyl$, or R^8 is halogen or NH_2 , and n is 0, 1, 2 or 3, preferably 0.

Preferred compounds of the formula (I) are those in which R^1 is H or (C_1-C_4) alkyl, preferably (C_1-C_4) alkyl,

R^2 and R^3 are H,

R^4 and R^5 are H,

R^6 is H,

R^7 is (C_1-C_4) alkyl, and

n is 0.

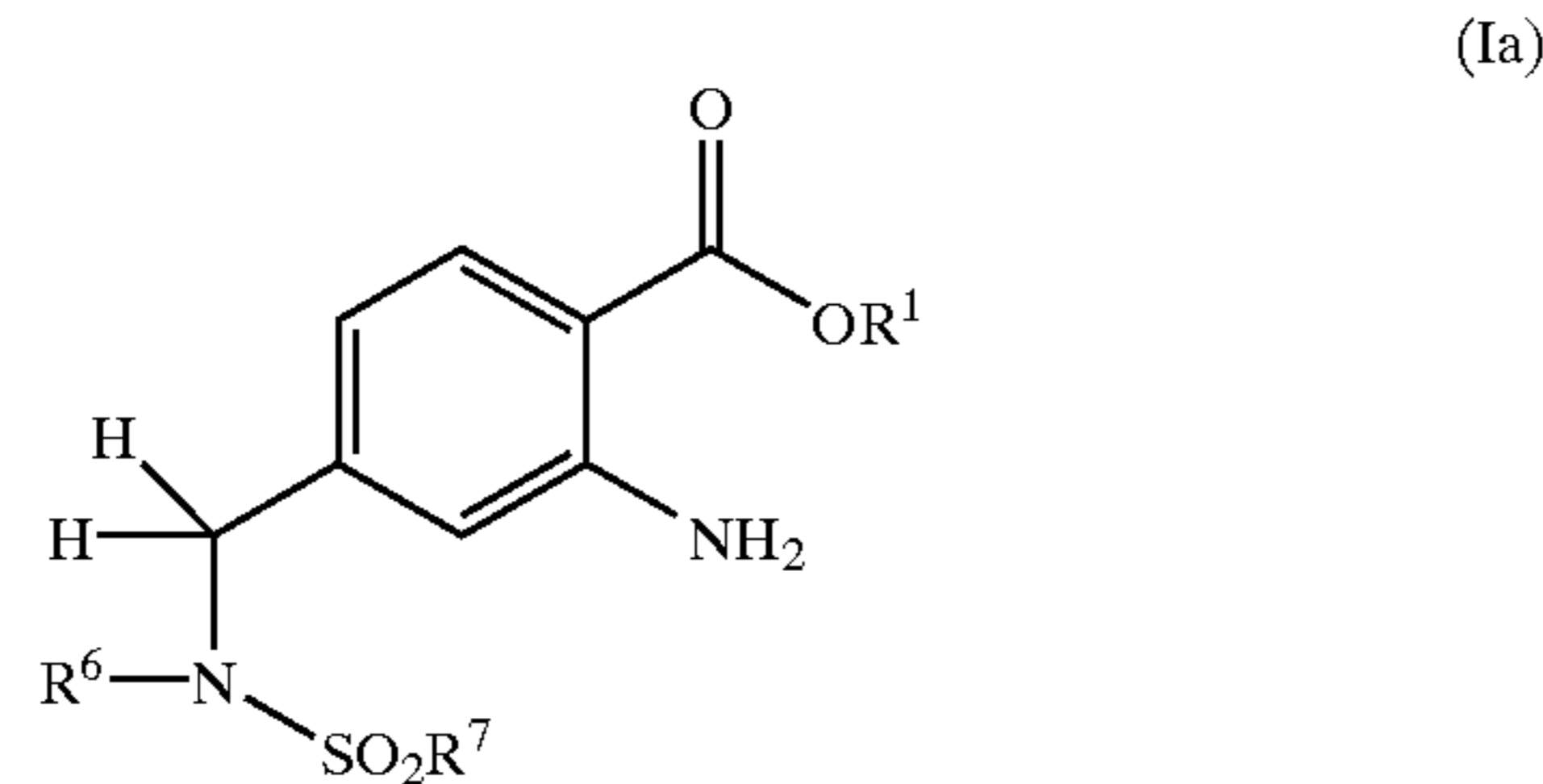
Compounds of the formula (I) which are of particular importance are those in which the group $CR^4R^5-NR^6-SO_2-R^7$ is in the para position relative to the group $-CO-OR^1$. If R^6 and R^7 together form a chain of the formula $-(CH_2)_m B_{m1}-$ and $m^1=1$, it is preferable that B is bound to the nitrogen atom which has R^6 attached to it.

Examples of compounds of the formula (I) are listed in table 1 hereinbelow:

TABLE 1

Compound	R^1	R^6	R^7
1	Me	H	Me
2	Me	Me	Me
3	Me	H	NHMe
4	Me	Me	NHMe
5	Me	H	N(Me) ₂
6	Me	Me	N(Me) ₂
7	Me	$-CH_2-CH_2-CH_2-$	
8	Me	$-CH_2-CH_2-CH_2-CH_2-$	
9	Me	H	Phe
10	Me	Me	Phe
11	Me	H	CH ₂ F
12	Me	Me	CH ₂ F
13	Me	H	CF ₃
14	Me	Me	CF ₃
15	Me	H	Et
16	Me	Me	Et
17	Me	H	nPr
18	Me	Me	nPr
19	Me	H	iPr
20	Me	Me	iPr
21	Me	H	nBu
22	Me	Me	nBu
23	Me	Et	Me
24	Me	Et	Et
25	Me	Et	NHMe
26	Me	Et	N(Me) ₂
27	Me	Et	Phe
28	Me	Et	CH ₂ F
29	Me	Et	CF ₃
30	Me	Et	nPr
31	Me	Et	iPr
32	Me	Et	nBu

TABLE 1-continued



(Ia)

Compound	R ¹	R ⁶	R ⁷
33	Et	H	Me
34	Et	Me	Me
35	Et	H	NHMe
36	Et	Me	NHMe
37	Et	H	N(Me) ₂
38	Et	Me	N(Me) ₂
39	Et	—CH ₂ —CH ₂ —CH ₂ —	
40	Et	—CH ₂ —CH ₂ —CH ₂ —CH ₂ —	
41	Et	H	Phe
42	Et	Me	Phe
43	Et	H	CH ₂ F
44	Et	Me	CH ₂ F
45	Et	H	CF ₃
46	Et	Me	CF ₃
47	Et	H	Et
48	Et	Me	Et
49	Et	H	nPr
50	Et	Me	nPr
51	Et	H	iPr
52	Et	Me	iPr
53	Et	H	nBu
54	Et	Me	nBu
55	Et	Et	Me
56	Et	Et	Et
57	Et	Et	NHMe
58	Et	Et	N(Me) ₂
59	Et	Et	Phe
60	Et	Et	CH ₂ F
61	Et	Et	CF ₃
62	Et	Et	nPr
63	Et	Et	iPr
64	Et	Et	nBu

In table 1, Me = methyl, Et = ethyl, nPr = n-propyl, iPr = isopropyl, nBu = n-butyl, Phe = phenyl.

If the term acyl is used in the present description, it denotes the radical of an organic acid which arises formally by eliminating an OH group from the organic acid, for example the radical of a carboxylic acid and radicals of acids derived therefrom, such as thiocarboxylic acid, optionally N-substituted iminocarboxylic acids or the radicals of carbonic monoesters, optionally N-substituted carbamic acids, sulfonic acids, sulfinic acids, phosphonic acids, phosphinic acids.

An acyl radical is preferably formyl or acyl from the group consisting of CO—R^x, CS—R^x, CO—OR^x, CS—OR^x, CS—SR^x, CR^x=NR^y, SOR^y or SO₂R^y, where R^x and R^y are each a C₁–C₁₀-hydrocarbon radical such as C₁–C₁₀-alkyl or C₆–C₁₀-aryl, each of which is unsubstituted or substituted, for example by one or more substituents selected from the group consisting of halogen such as F, Cl, Br, I, alkoxy, haloalkoxy, hydroxyl, amino, nitro, cyano or alkylthio, or acyl is aminocarbonyl or aminosulfonyl, the two last-mentioned radicals being unsubstituted, N-monosubstituted or N,N-disubstituted, for example by substituents from the group consisting of alkyl or aryl. Acyl is, for example, formyl, haloalkylcarbonyl, alkylcarbonyl such as (C₁–C₄)alkylcarbonyl, phenylcarbonyl, it being pos-

sible for the phenyl ring to be substituted, or alkyloxycarbonyl, such as (C₁–C₄)alkyloxycarbonyl, phenyloxycarbonyl, benzyloxycarbonyl, alkylsulfonyl, such as (C₁–C₄)alkylsulfonyl, alkylsulfinyl, such as C₁–C₄ (alkylsulfinyl), N-alkyl-1-iminoalkyl, such as N-(C₁–C₄)-1-imino-(C₁–C₄)alkyl and other radicals of organic acids.

In formula (I) and the general formulae used hereinbelow, the radicals alkyl, alkoxy, haloalkyl, haloalkoxy and alkylthio and the corresponding substituted radicals can be in each case straight-chain or branched in the carbon skeleton. Unless specified otherwise, the lower carbon skeletons, for example those having 1 to 4 carbon atoms, are preferred amongst these radicals. Alkyl radicals, also in the composite meanings such as alkoxy, haloalkyl and the like, are, for example, methyl, ethyl, n- or i-propyl, n-, i-, t- or 2-butyl, pentyls, hexyls such as n-hexyl, i-hexyl and 1,3-dimethylbutyl, heptyls such as n-heptyls, 1-methylhexyl and 1,4-dimethylpentyl; alkenyl and alkynyl radicals have the meanings of the possible unsaturated radicals which correspond to the alkyl radicals; for example alkenyl is allyl, 1-methylprop-2-en-1-yl, 2-methylprop-2-en-1-yl, but-2-en-1-yl, but-3-en-1-yl, 1-methylbut-3-en-1-yl and 1-methylbut-2-en-1-yl; alkynyl is, for example, propargyl, but-2-yn-1-yl, but-3-yn-1-yl, 1-methylbut-3-yn-1-yl.

Alkenyl, for example in the form “(C₃–C₈)alkenyl”, is preferably an alkenyl radical having 3 to 8 carbon atoms in which the double bond is not positioned at the carbon atom which is linked to the remaining moiety of the compound (I) (“yl” position). This also applies analogously to alkynyl radicals.

Halogen is, for example, fluorine, chlorine, bromine or iodine. Haloalkyl, -alkenyl and -alkynyl are alkyl, alkenyl or alkynyl, each of which is partially or fully substituted by halogen, preferably by fluorine, chlorine and/or bromine, in particular by fluorine or chlorine, for example CF₃, CHF₂, CH₂F, CF₃CF₂, CH₂FCHCl₂, CCl₃, CHCl₂, CH₂CH₂Cl; haloalkoxy is, for example, OCF₃, OCHF₂, OCH₂F, CF₃CF₂O, OCH₂CF₃ and OCH₂CH₂Cl; this also applies analogously to haloalkenyloxy and other halogen-substituted radicals.

Substituted radicals such as substituted hydrocarbon radicals, for example substituted alkyl, alkenyl, alkynyl, aryl, for example phenyl, are, for example, a substituted radical which is derived from the unsubstituted skeleton, the substituents being, for example, one or more, preferably 1, 2 or 3, radicals selected from the group consisting of halogen, alkoxy, haloalkoxy, alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido, alkyloxycarbonyl, alkylcarbonyl, formyl, carbamoyl, mono- and dialkylaminocarbonyl, substituted amino such as acylamino, mono- and dialkylamino, and alkylsulfinyl, haloalkylsulfinyl, alkylsulfonyl, haloalkylsulfonyl and, in the case of cyclic radicals, also alkyl and haloalkyl, and unsaturated aliphatic radicals which correspond to the abovementioned saturated hydrocarbon-containing radicals, such as alkenyl, alkynyl, alkenyloxy, alkynyloxy and the like. In the case of radicals with carbon atoms, those having 1 to 4 carbon atoms, in particular 1 or 2 carbon atoms, are preferred. Preferred are, as a rule, substituents selected from the group consisting of halogen, for example fluorine and chlorine, (C₁–C₄)alkyl, preferably methyl or ethyl, (C₁–C₄)

5

haloalkyl, preferably trifluoromethyl, (C₁-C₄)alkoxy, preferably methoxy or ethoxy, (C₁-C₄)haloalkoxy, nitro and cyano. Especially preferred in this context are substituents methyl, methoxy and chlorine.

Optionally substituted phenyl or phenoxy is preferably phenyl or phenoxy, each of which is unsubstituted or mono- or polysubstituted, preferably up to trisubstituted, by identical or different radicals selected from the group consisting of halogen, (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄) haloalkyl, (C₁-C₄)haloalkoxy and nitro, for example o-, m- and p-tolyl, dimethylphenyls, 2-, 3- and 4-chlorophenyl, 2-, 3- and 4-trifluoro- and -trichlorophenyl, 2,4-, 3,5-, 2,5- and 2,3-dichlorophenyl, o-, m- and p-methoxyphenyl.

If substitutions are defined by one or more radicals from among a group of radicals, this encompasses both the substitution by one or more identical radicals and the mono- or polysubstitution by different radicals.

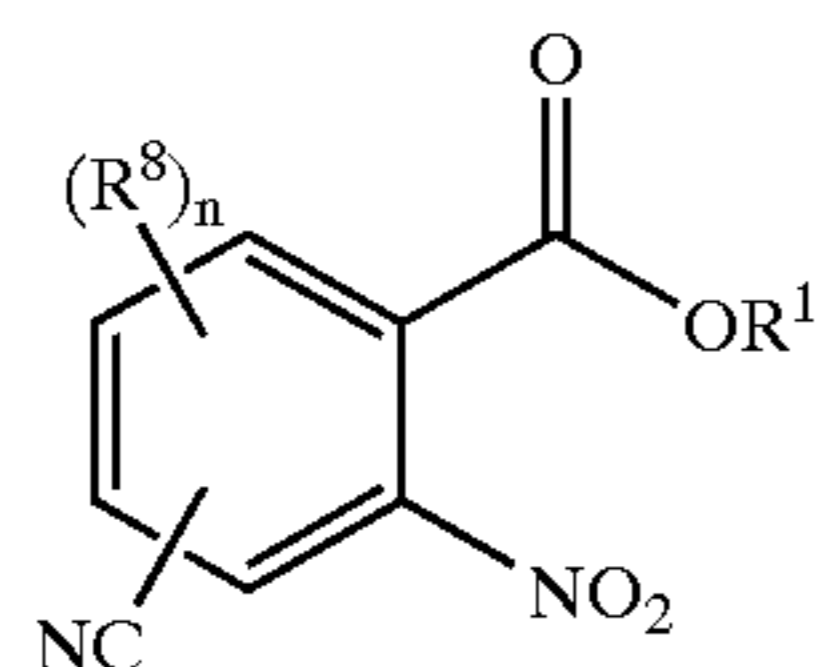
Subject of the invention are also all stereoisomers which are encompassed by formula (I) and their mixtures. Such compounds of the formula (I) contain one or more asymmetric carbon atoms which are not indicated separately in formula (I). The possible stereoisomers which are defined by their specific spatial shape, such as enantiomers or diastereomers, are all encompassed by formula (I) and can be obtained from mixtures of the stereoisomers by customary methods or else by stereoselective reactions in combination with the use of stereochemically pure starting materials. Formula (I) also encompasses tautomers of the compounds stated, insofar as they are formed by proton migration and are chemically stable.

The compounds of the formula (I) may form salts in which an acidic hydrogen atom is replaced by a suitable cation. These salts are, for example, metal salts; preferably alkali metal salts or alkaline earth metal salts, in particular sodium salts and potassium salts, or else ammonium salts or salts of organic amines. Likewise, salt formation can be effected by an addition reaction of an acid with basic groups, such as amino. Acids which are suitable for this purpose are strong inorganic and organic acids, for example HCl, HBr, H₂SO₄, HNO₃ or formic acid.

Compounds of the formula (I) are successfully synthesized in very good yields and purities starting from compounds of the formula (II) mentioned hereinbelow.

Subject of the present invention is thus also a process for the preparation of compounds of the formula (I) comprising the steps of

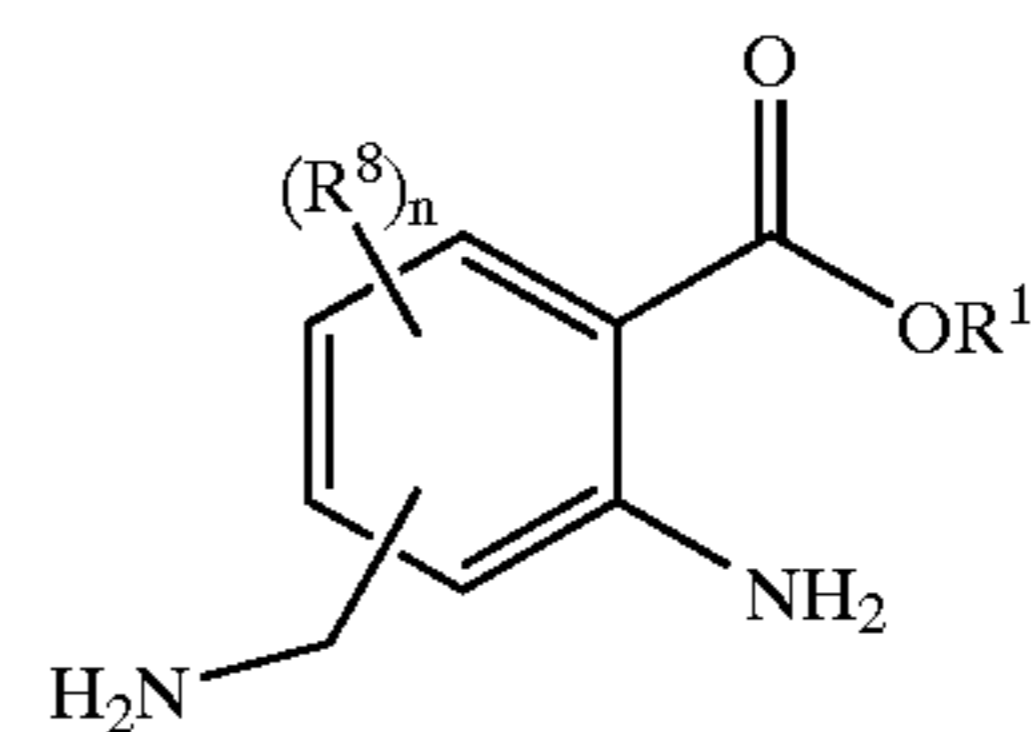
1a) reacting a compound of the formula (II)



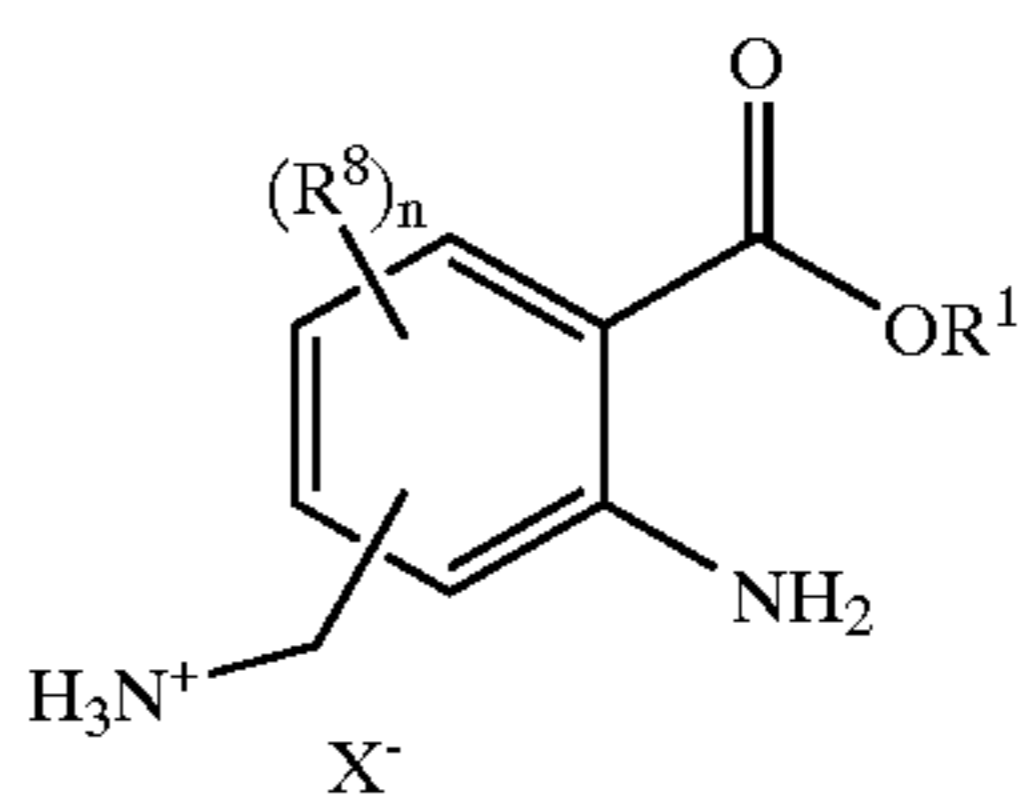
by catalytic hydrogenation in the absence of an acid to give a compound of the formula (III) or by catalytic hydrogenation in the presence of an acid, for example H⁺X⁻, where X⁻ is an equivalent of an acid anion, such as

6

halogen, for example Cl⁻, Br⁻ or I⁻, or HSO₄⁻, ½ SO₄²⁻, H₂PO₄⁻, ½ HPO₄²⁻, ⅓ PO₄³⁻ or ⁻OCOR (where R=H or (C₁-C₈)alkyl) to give a compound of the formula (IIIa), where X⁻ is an equivalent of an acid anion, such as halide, for example Cl⁻, Br⁻ or I⁻, or HSO₄⁻, ½ SO₄²⁻, H₂PO₄⁻, ½ HPO₄²⁻, ⅓ PO₄³⁻ or ⁻OCOR (where R=H or (C₁-C₈)alkyl),



(III)

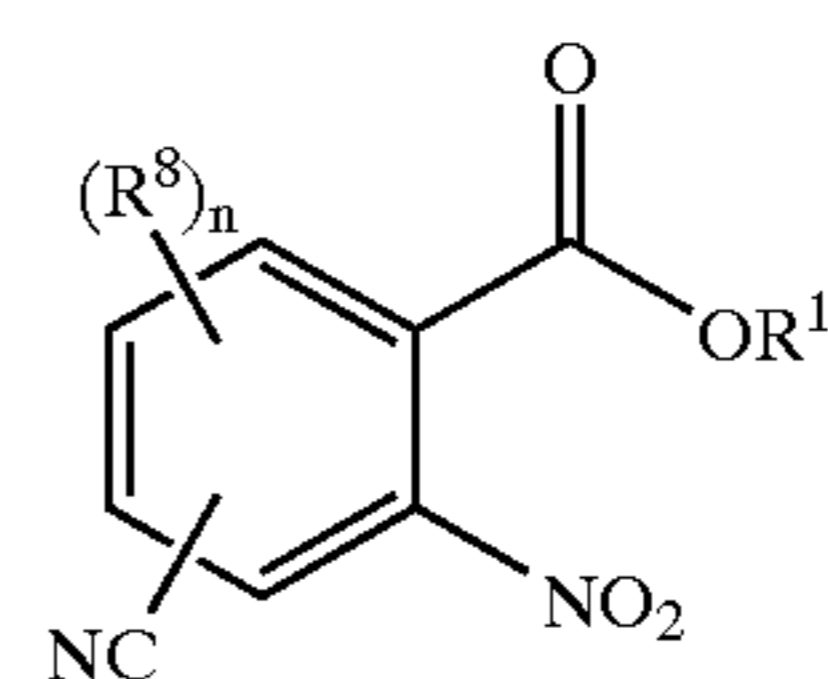


(IIIa)

and subsequently

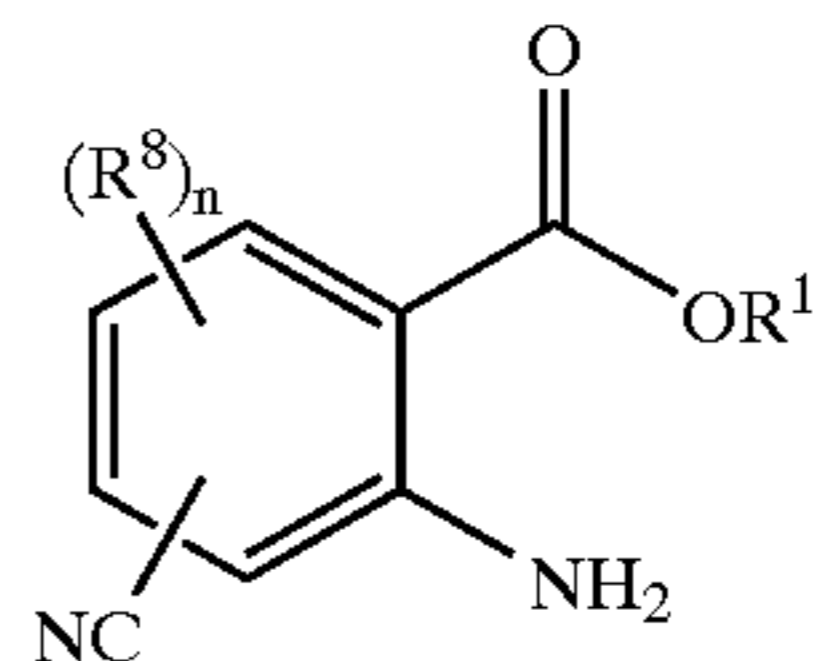
1b) reacting the compound of the formula (III) or (IIIa) with a sulfonic acid derivative to give a compound of the formula (I) where R², R³ and R⁶=H; or

2a) α) reacting a compound of the formula (II)



(II)

by customary reduction methods for nitro compounds to give a compound of the formula (IV),



(IV)

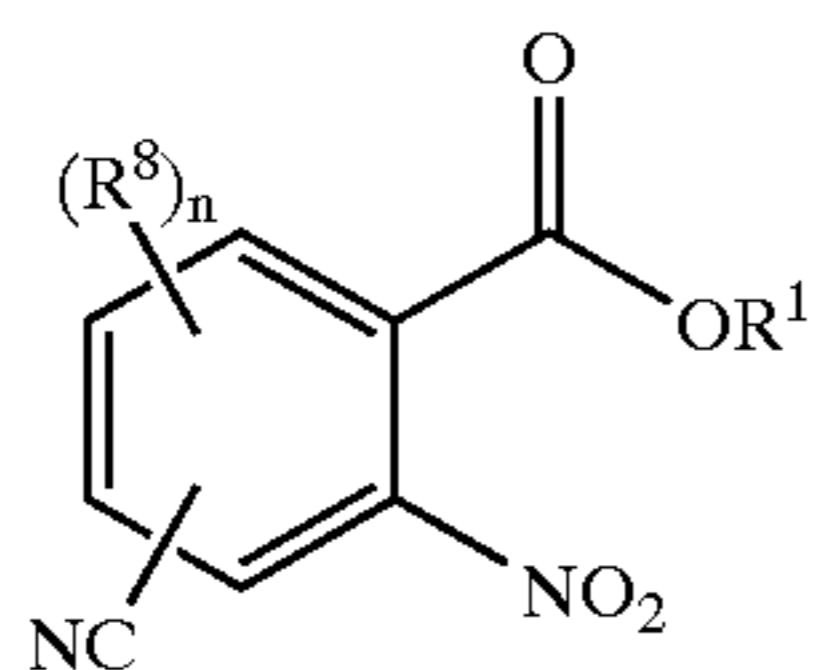
and subsequently

β) reacting the compound of the formula (IV) either by catalytic hydrogenation or by customary reduction methods for nitriles to give a compound of the formula (III) or (IIIa),

and subsequently

2b) reacting the compound of the formula (III) or (IIIa) with a sulfonic acid derivative to give a compound of the formula (I) where R², R³ and R⁶=H; or

7

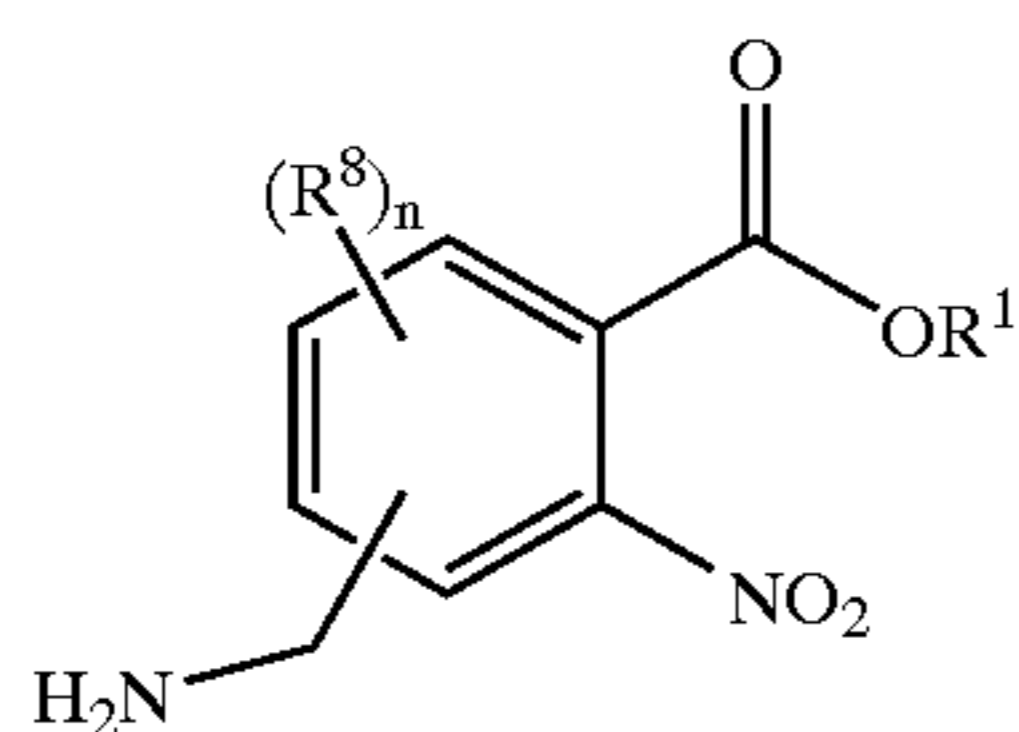
3a) α) reacting a compound of the formula (II)

(II)

5

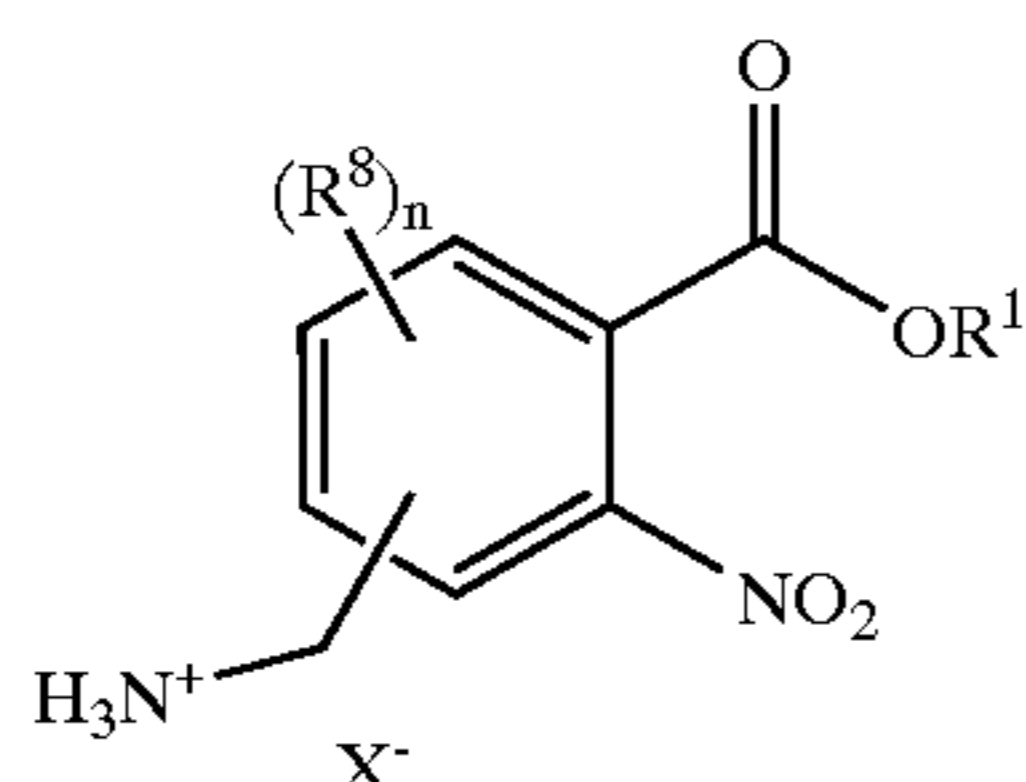
10

by customary reduction methods for nitrites to give a compound of the formula (V) or (Va), where X^{\ominus} is as defined in formula (IIIa),



(V)

20



(Va)

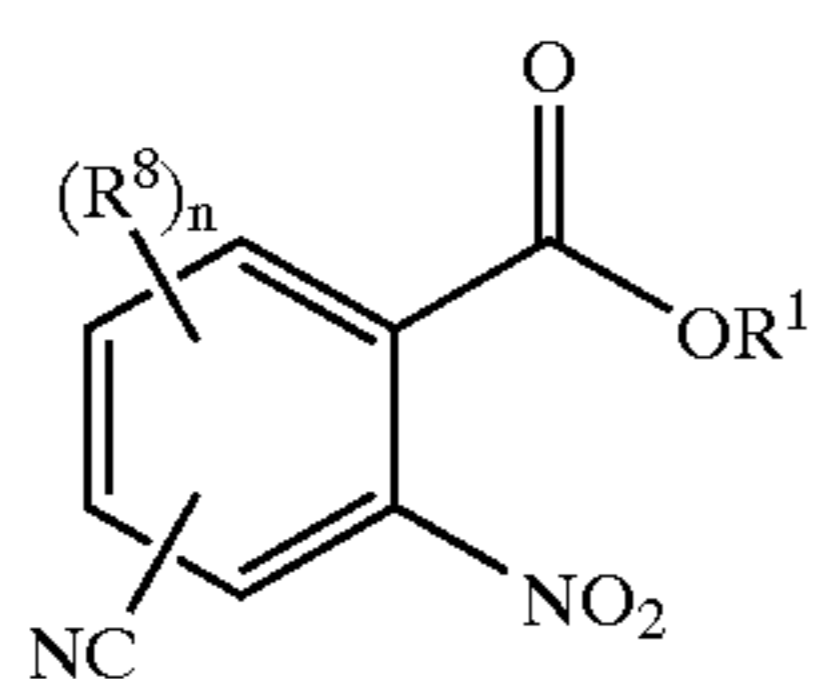
25

and subsequently

β) reacting the compound of the formula (V) or (Va) by customary reduction methods for nitro compounds or by catalytic hydrogenation to give a compound of the formula (III) or (IIIa),

and subsequently

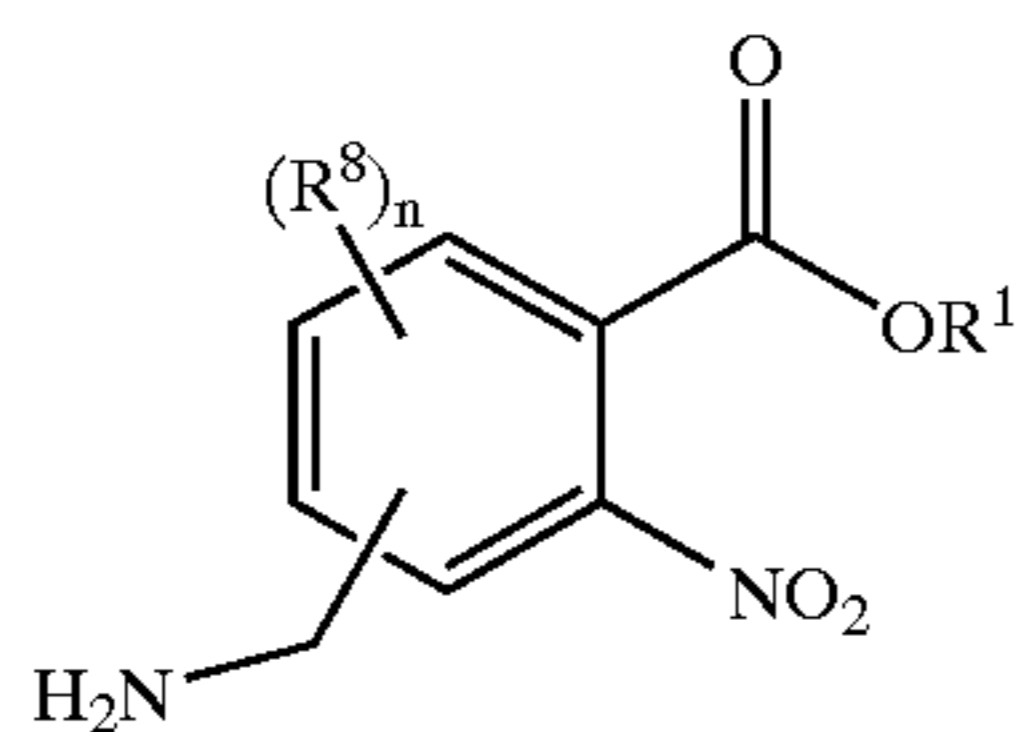
3b) reacting the compound of the formula (III) or (IIIa) with a sulfonic acid derivative to give a compound of the formula (I) where R^2 , R^3 and $R^6=H$; or

4a) α) reacting a compound of the formula (II)

(II)

50

by customary reduction methods for nitrites to give a compound of the formula (V) or (Va), where X^- is as defined in formula (IIIa),



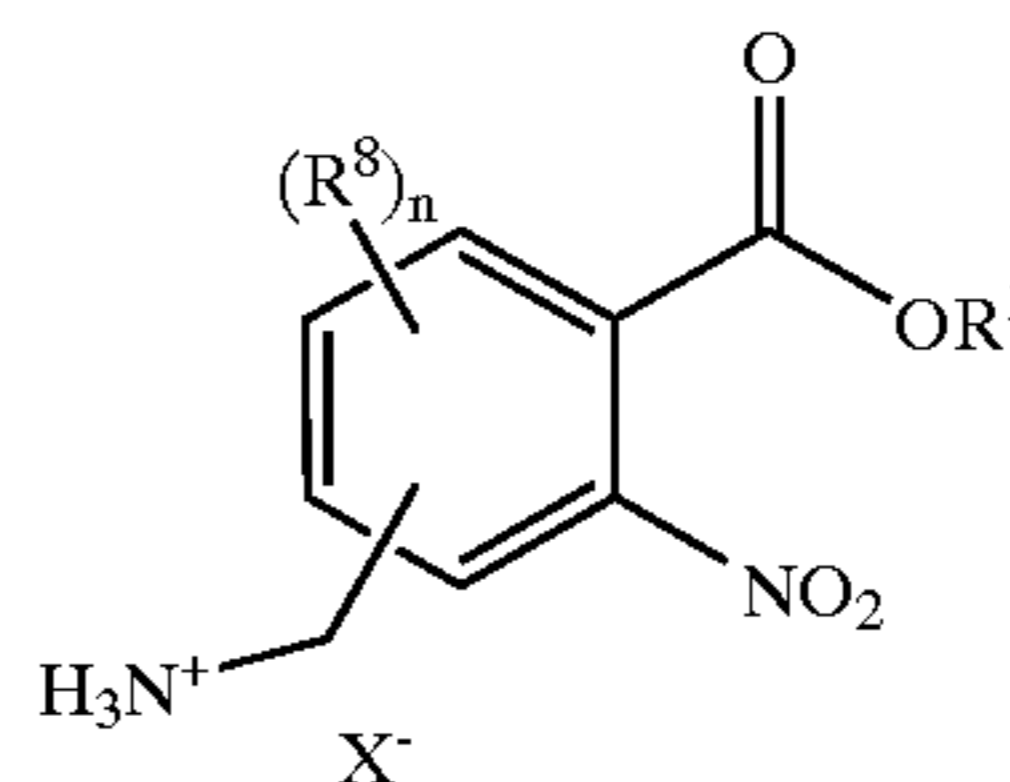
(V)

65

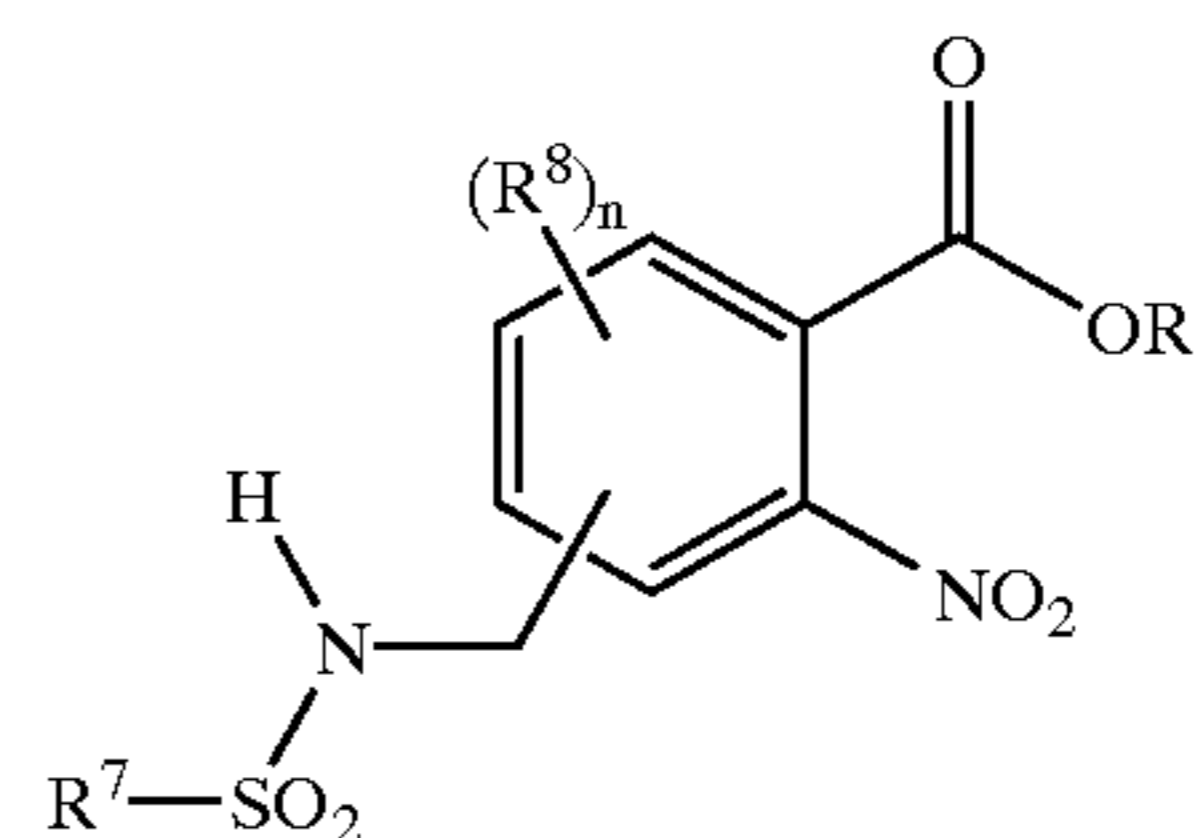
8

-continued

(Va)



β) and subsequently reacting the compound of the formula (V) or (Va) with a sulfonic acid derivative to give a compound of the formula (VI),



(VI)

and subsequently

4b) reacting the compound of the formula (VI) by customary reduction methods for nitro compounds or by catalytic hydrogenation to give a compound of the formula (I) where R^2 , R^3 and $R^6=H$.

The compounds of formulae (III), (IIIa), (V), (Va) and (VI) are novel and also subject of the present invention.

Compounds of the formula (I) where R^2 and/or $R^3=acyl$ can be obtained by acylating compounds of the formula (I) where R^2 and $R^3=H$ with acylating agents such as carbonyl halides, sulfonyl halides and carbamoyl halides, carboxylic anhydrides, sulfonic anhydrides, haloformic esters or isocyanates, by customary methods (see, for example, L.-F. Tietze, Th. Eicher, *Reaktionen und Synthesen im organisch-chemischen Praktikum [Reactions and Syntheses in the Organochemical Laboratory Practical]*, Thieme Verlag Stuttgart/New York, 1981, pp. 131, 316, 318, 345; R. C. Larock, *Comprehensive Organic Transformations* (1989), pp. 979, 981). Examples of suitable solvents are aprotic solvents such as dichloromethane, acetonitrile, dioxane, tetrahydrofuran, toluene or chlorobenzene, preferably at temperatures of from 0° C. to the boiling point of the solvent.

Compounds of the formula (I) where $R^6=unsubstituted$ or substituted C_1-C_8 -alkyl can be obtained for example by alkylating compounds of the formula (I) where $R^6=H$ with alkylating agents such as alkyl halides, alkyl sulfates such as dimethyl sulfate or alkyl tosylates by customary methods. Examples of suitable solvents are acetone and dimethylformamide (cf., for example, R. C. Larock, *Comprehensive Organic Transformations* (1989), p. 398; L.-F. Tietze, Th. Eicher, *Reaktionen und Synthesen im organisch-chemischen Praktikum*, Thieme Verlag Stuttgart/New York, 1981, p. 75; *Organikum, Organisch-chemisches Grundpraktikum [Basic Laboratory Practical in Organic Chemistry]* VEB, Berlin 1981). The alkylation can be carried out in the presence of bases such as K_2CO_3 , NaH or alkoxides such as sodium alkoxide. The starting material is preferably compounds of the formula (I) in which R^2 and R^3 are acyl.

Compounds of the formula (I) where R⁶=unsubstituted or substituted C₁-C₈-alkyl can also be obtained for example via reductive aminations, for example with aldehydes or ketones in the presence of reducing agents such as H₂/catalyst, formic acid, zinc/HCl, sodium borohydride or sodium cyanoborohydride. An example is the Leuckard-Wallach reaction with formaldehyde and formic acid.

Preferred processes are those in which the nitro and the nitrile group in compounds of the formula (II) are reduced jointly in one process step by means of catalytic hydrogenation in accordance with process variant 1a) to give compounds of the formula (III) or (IIIa).

The amino compounds of the formulae (III) and (V) which are obtained as intermediates in the process according to the invention can also arise in the form of their salts (IIIa) and (Va) and can be reacted further when the reaction or work-up is effected in an acidic medium.

The symbols given in formulae (II), (III), (IIIa), (IV), (V), (Va) and (VI) have the same meaning as in formula (I), including the preferred ranges mentioned herefor. Preferred compounds of the formulae (II), (III), (IIIa), (IV), (V), (Va) and (VI) are those in which the groups —CN (formulae (II) and (IV)), —CH₂—NH₂ (formulae (III) and (V)), —CH₂—NH₃⁺X⁻ (formulae (IIIa) and (Va)) and —CH₂—SO₂—R⁷ (formula (VI)) are in the para position relative to the group —CO—OR¹.

Furthermore, substeps of the process according to the invention are also subject of the invention.

The compounds of the formula (II) are known, cf., for example, DE 22 39 799 C3 or Journal of the American Chemical Society 99, 6721 (1977).

The catalytic hydrogenation of the compound of the formula (II) by process variant 1a), of the compound of the formula (IV) by process variant 2aβ), of the compound of the formula (V) or (Va) by process variant 3aβ) or of the compound of the formula (VI) by process variant 4b) is successfully carried out by means of customary hydrogenation methods. Examples of hydrogen sources which can be used are hydrogen gas, hydrazine or HN=NH. Particularly suitable hydrogenation catalysts are noble-metal catalysts, for example Pd, Pt, Rh, Ir or Ni or Co catalysts. The noble metals can be used in elemental form or in the form of oxides or halides. The noble-metal catalysts can be used as desired without or, preferably, with support materials such as active charcoal, kieselguhr, silicates.

The hydrogenation can be carried out both by atmospheric pressure and by applying a superatmospheric hydrogen pressure, as a rule between 1 and 100 bar, preferably 1–50 bar. In general, the suitable temperature is in the range of from –20 to 150° C., preferably between 0 and 120° C.

Examples of solvents which are suitable for the hydrogenation are solvents of the groups water, alcohols such as methanol or ethanol, ethers such as diethyl ether, tetrahydrofuran or dioxane, amides such as dimethylformamide or dimethylacetamide, esters such as ethyl acetate, organic carboxylic acids such as formic acid or acetic acid, aromatic hydrocarbons such as toluene, xylene and chlorobenzene, or halogenated aliphatic hydrocarbons such as CH₂Cl₂, it being possible to employ the solvents in pure form or as mixtures.

The catalytic hydrogenation of the compounds of the formula (II) by process variant 1a) or of the compounds of

the formula (IV) by process variant 2aβ) is preferably carried out in the presence of 1–10 molar equivalents of an acid. Solvents which are preferably used are alcohols such as methanol or ethanol, or water. Examples of suitable acids are inorganic acids or carboxylic acids. Preferred are acids of the formula H⁺X⁻ where X⁻ is an equivalent of an acid moiety, such as halogen for example Cl⁻, Br⁻ or I⁻, or HSO₄⁻, ½ SO₄²⁻, H₂PO₄⁻, ½ HPO₄²⁻, ⅓ PO₄³⁻ or -OCOR (where R=H or (C₁-C₈)alkyl), for example hydrohalic acids such as hydrochloric acid or hydrobromic acid, or sulfuric acid, phosphoric acid, formic acid or acetic acid. If, for example, the two last-mentioned acids are used, the acids may also fully assume the role of the solvent.

The catalytic hydrogenation of the compounds of the formula (IV) can also be carried out by using 1–10 molar equivalents of ammonia, nickel or cobalt catalysts such as Raney nickel or Raney cobalt preferably being employed. Solvents which are preferably used in this context are alcohols such as methanol or ethanol.

The reduction of the nitro group in compounds of the formula (II) by process variant 2aα), compounds of the formulae (V) and (Va) by process variant 3aβ) or compounds of the formula (VI) by process variant 4b) can be carried out with customary reducing agents for aromatic nitro compounds. Such reducing agents and reaction conditions are described, for example, in R. C. Larock, Comprehensive Organic Transformations (1989) pp. 411–415, VCH Publishers Inc. and the literature cited therein. Examples of preferred reducing agents are Fe, Zn, Sn or their salts such as FeSO₄ or Sn-II salts such as SnCl₂. Examples of suitable solvents are organic carboxylic acids, alcohols and/or mineral acids. In general, the reaction temperature is between 0° C. and the boiling point of the solvent.

The reduction of the nitrile group in compounds of the formula (IV) by process variant 2aβ) and compounds of the formula (II) by process variant 3aα) and 4aα) can be carried out by customary reducing agents for nitriles. Such reducing agents and reaction conditions are described, for example, in R. C. Larock, Comprehensive Organic Transformation (1989) pp. 437–438, VCH Publishers Inc. and the literature cited therein. Examples of preferred reducing agents are boron hydride compounds or aluminum hydride compounds such as BH₃/THF, BH₃/DMS and their salts such as NaBH₄. Examples of suitable solvents are ethers such as dioxane or tetrahydrofuran. The reaction temperature is generally between 0° C. and the boiling point of the solvent. If the reduction product is subsequently worked up in an acid medium, for example methanol/HCl, the compound of the formula (III) (process variant 2aβ) or the compound of the formula (V) (process variants 3aα and 4aα) can be obtained in the form of a salt of the formula (IIIa) or (Va), respectively, which can be reacted further analogously to compound (III) or compound (V), respectively.

The acylation of the compounds of the formula (III) or (IIIa) by process variant 1b), 2b) or 3b) or of the compounds of the formula (V) or (Va) by process variant 4aβ) with a sulfonic acid derivative can be carried out under customary conditions for acylation reactions to give the compounds of the formula (VI) in the case of compounds of the formula (V) or (Va) or to give the compounds of the formula (I) according to the invention in the case of compounds of the formula (III) or (IIIa).

11

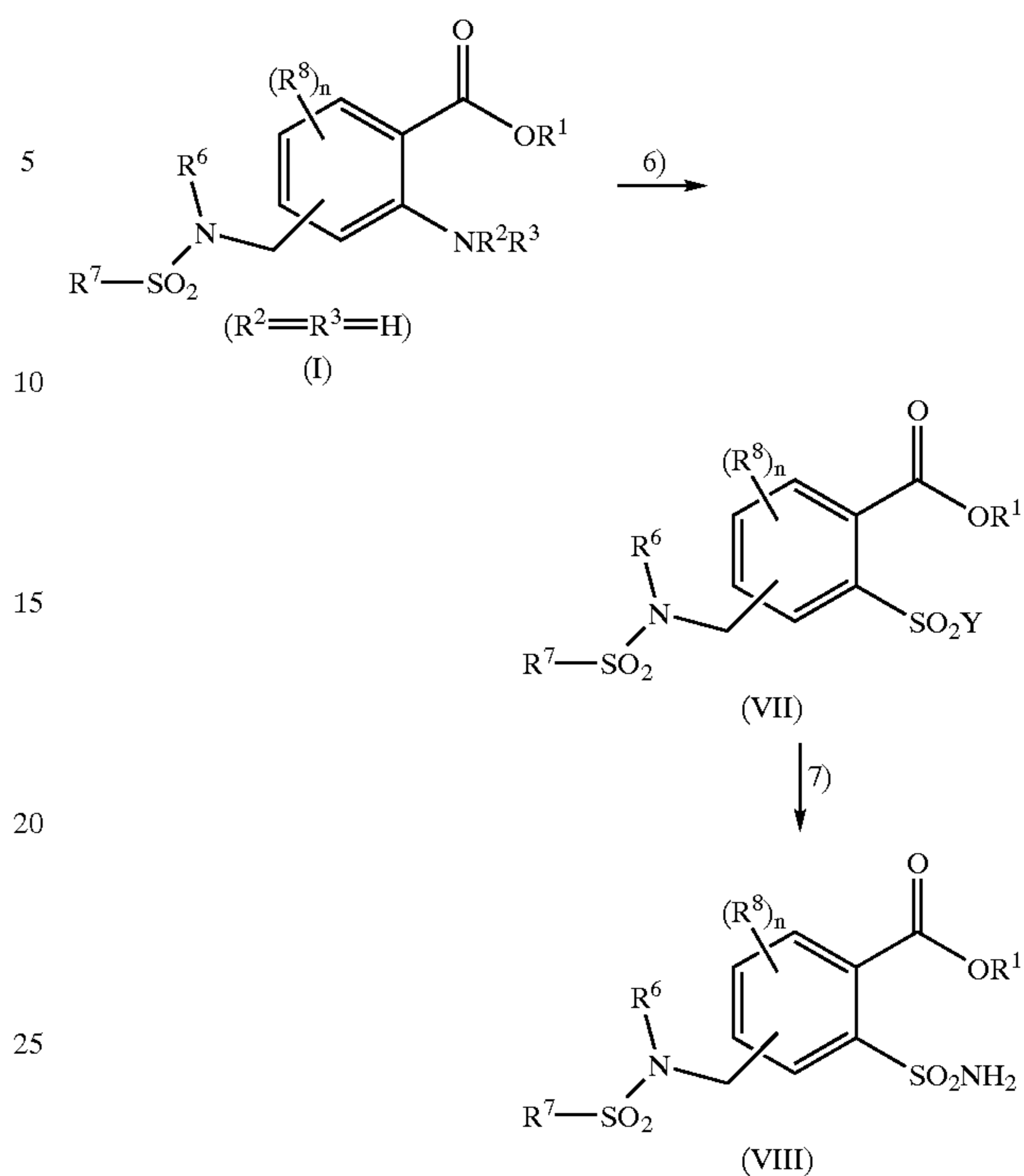
For example, compounds of the formula (III) or (IIIa), or (V) or (Va), are reacted in suitable solvents with sulfonic acid derivatives in the presence of bases as acid acceptors to give compounds of the formula (I) or (VI) respectively. Examples of solvents which are suitable for the acylations are solvents from the groups water, alcohols such as methanol or ethanol, halogenated aliphatic hydrocarbons such as CH_2Cl_2 , aromatic hydrocarbons such as toluene, chlorobenzene or xylene, ethers such as diethyl ether, tetrahydrofuran or dioxane, ketones such as acetone or methyl isobutyl ketone, esters such as ethyl acetate, and aprotic solvents such as acetonitrile, dimethylformamide or dimethylacetamide, it being possible for the solvents to be employed in pure form or as mixtures. Preferred are water and mixtures of water and water-soluble organic solvents from the abovementioned groups.

Bases which are suitable are inorganic or organic bases, for example carbonates such as K_2CO_3 , Na_2CO_3 or NaHCO_3 , alkali metal hydroxides and alkaline earth metal hydroxides such as NaOH , KOH or $\text{Ca}(\text{OH})_2$, or amines such as triethylamine. In general, the bases are employed in amounts of 1–10 molar equivalents, preferably 1–5 molar equivalents, per compound of the formula (III) or (V); when compounds of the formula (IIIa) or (Va) are employed, the minimum amount of the base employed is at least two molar equivalents.

Examples of suitable sulfonic acid derivatives are sulfonyl halides such as fluorides, chlorides, bromides or iodides, and sulfonic anhydrides. Preferred are sulfonic acid derivatives of the formula $\text{R}^7\text{—SO}_2\text{—Z}$, where R^7 is defined as in formula (I), and Z is a leaving group such as halogen (for example fluorine, chlorine, bromine or iodine) or $\text{O—SO}_2\text{—R}^Z$, where R^Z is as defined for R^7 in formula (I). The acylation is carried out for example in such a way that the compounds of the formula (III) or (IIIa), or (V) or (Va), are reacted with the sulfonic acid derivatives in suitable solvents in the presence of a suitable base, in general at temperatures of from -20 to 100°C . Preferred are temperatures of from -10 to 50°C . The amounts of sulfonic acid derivatives are generally 1–10 molar equivalents, preferably 1–5 molar equivalents, per compound of the formula (III) or (IIIa), or (V) or (Va).

In addition to compounds of the formula (I) and their preparation, the present invention also relates to their further reaction to give compounds of the formulae (VII) and (VIII). To do this, compounds of the formula (I) where R^2 and/or $\text{R}^3=\text{acyl}$ must first be converted by customary methods into compounds of the formula (I) where $\text{R}^2=\text{R}^3=\text{H}$, and these are then further reacted to give compounds of the formulae (VII) and (VIII). The symbols used in formulae (VII) and (VIII) have the same meanings as stated for formula (I), including the preferred ranges stated herefor, and Y in formula (VII) is halogen such as fluorine, chlorine, bromine or iodine. Preferred compounds of the formulae (VII) and (VIII) are those in which the group $\text{—CH}_2\text{—NR}^6\text{—SO}_2\text{—R}^7$ is in the para position relative to the group —CO—OR^1 .

12



As is described in (EP-A-723 534), compounds of the formulae (VII) and (VIII) are suitable precursors for the preparation of potent herbicidal sulfonylureas, the preparation of the compounds of the formulae (VII) and (VIII) being especially efficient in the present process according to the invention and the compounds of the formulae (VII) and (VIII) being obtained in very good yields and purities.

Methods for the conversion 6) of anilines into sulfonyl halides are known (see, for example, H. Meerwein et al., Chem. Berichte 90, 841–852 (1957)). Surprisingly, compounds of the formula (I) where $\text{R}^2, \text{R}^3=\text{H}$ are successfully reacted to give compounds of the formula (VII) on the basis of procedures described in the literature. Thus, compounds of the formula (I) where $\text{R}^2, \text{R}^3=\text{H}$ can be diazotized under suitable conditions and subsequently coupled with suitable SO_2 sources, such as SO_2 gas, $\text{Na}_2\text{S}_2\text{O}_5$ or NaHSO_3 in the presence of acids such as carboxylic acids, for example acetic acid, or inorganic acids, for example hydrohalic acids HY such as HCl or HBr , and catalysts, for example copper catalysts based on Cu(I) and/or Cu(II) salts to give sulfonyl halides of the formula (VII).

The diazotization can be carried out with suitable diazotizing agents such as NaNO_2 in the presence of acids such as inorganic acids, preferably hydrohalic acids HY, such as HCl or HBr . The solvent used is preferably a water/acid mixture, in particular a mixture of water/carboxylic acid (for example acetic acid) or water/mineral acid (for example hydrohalic acid HY such as HCl or HBr). In general, the reaction temperature is -20 to 50°C ., preferably -10 to 20°C .

The following are examples which can be used as solvents for the subsequent coupling reaction: water, carboxylic acids such as acetic acid, carboxylic esters such as ethyl acetate, ethers such as diethyl ether, tetrahydrofuran or dioxane, halogenated aliphatic hydrocarbons such as CH_2Cl_2 or

13

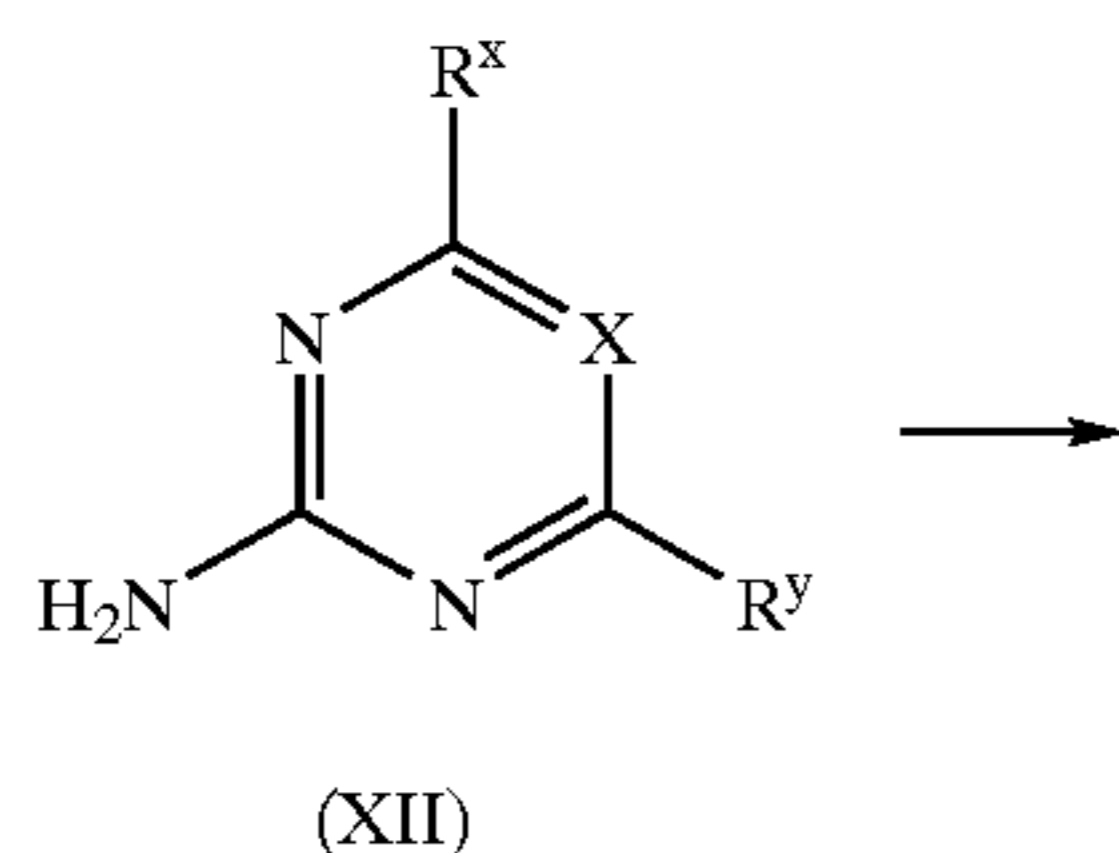
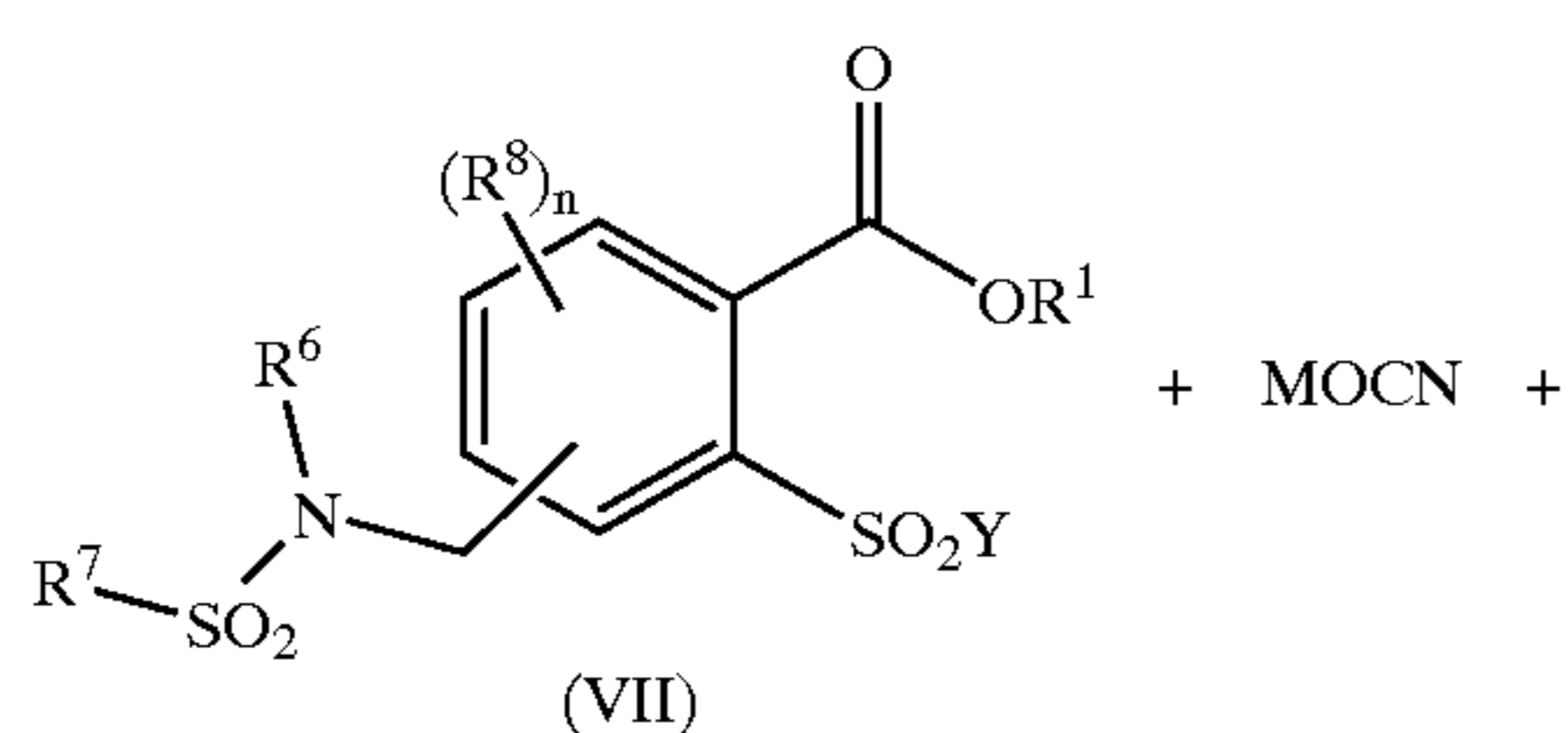
dichloroethane, aromatic hydrocarbons such as toluene, chlorobenzene or xylene, or ketones such as acetone or methyl isobutyl ketone. Moreover, the reaction mixture contains acids, for example carboxylic acids such as acetic acid or mineral acids such as hydrohalic acids HY, for example HCl or HBr, which are either still present from the diazotization reaction and/or are added when the coupling reaction is carried out. Examples of SO₂ sources which can be used are, for example, SO₂ gas (1–10 equivalents), Na₂S₂O₅ (1–10 equivalents) or NaHSO₃ (1–10 equivalents), in the presence of catalysts, for example copper catalysts such as CuCl (1–20 mol %), CuCl₂ (1–20 mol %), CuBr (1–20 mol %) or CuBr₂ (1–20 mol %).

Starting from sulfonyl halides of the formula (VII) the aminolysis 7) which yields sulfonic amides of the formula (VIII) is, surprisingly, successfully carried out with high efficiency and in high yields by reacting compounds of the formula (VII) for example in suitable solvents with ammonia.

The aminolysis can be carried out with suitable reagents, for example 2–10 molar equivalents of aqueous ammonia solution or NH₃ gas in the presence of a solvent, for example ketones such as acetone or methyl isobutyl ketone, halogenated aliphatic hydrocarbons such as CH₂Cl₂, aromatic hydrocarbons such as xylene, toluene or chlorobenzene, ethers such as diethyl ether, tetrahydrofuran or dioxane, esters such as ethyl acetate, aprotic solvents such as dimethylformamide, dimethylacetamide or acetonitrile, or mixtures of these solvents. In general, the reaction temperature is from –10 to 100° C., preferably –10 to 40° C., especially preferably –10 to 20° C.

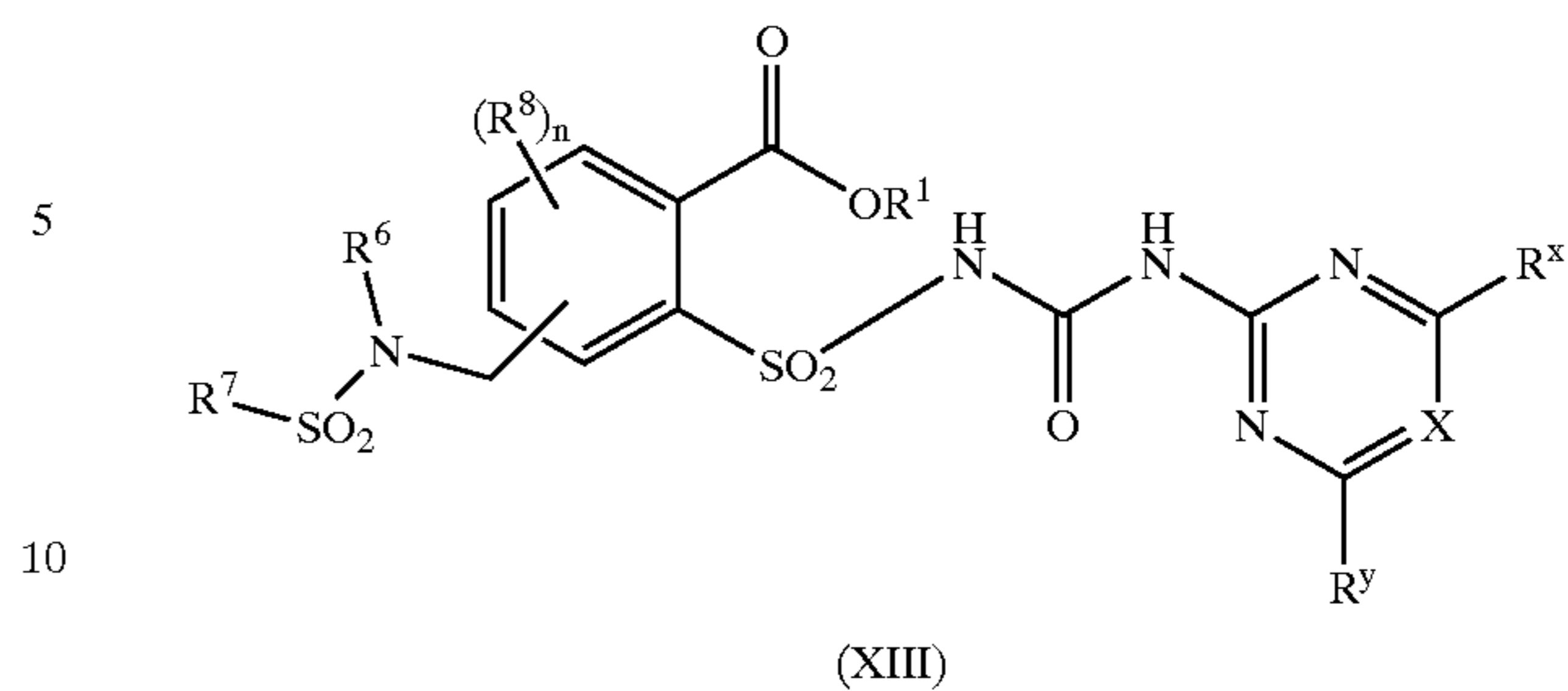
The compounds of the formulae (VII) and (VIII) can subsequently be reacted in various ways to give sulfonylureas, preferably sulfonylureas of the formula (XIII) and/or their salts, for example by

8) reacting a sulfonyl halide of the formula (VII) with a cyanate MOCN, in which M is an ammonium ion or an alkali metal ion such as Li, Na or K, and with an amino heterocycle of the formula (XII) in the presence of a base to give the sulfonylurea; or

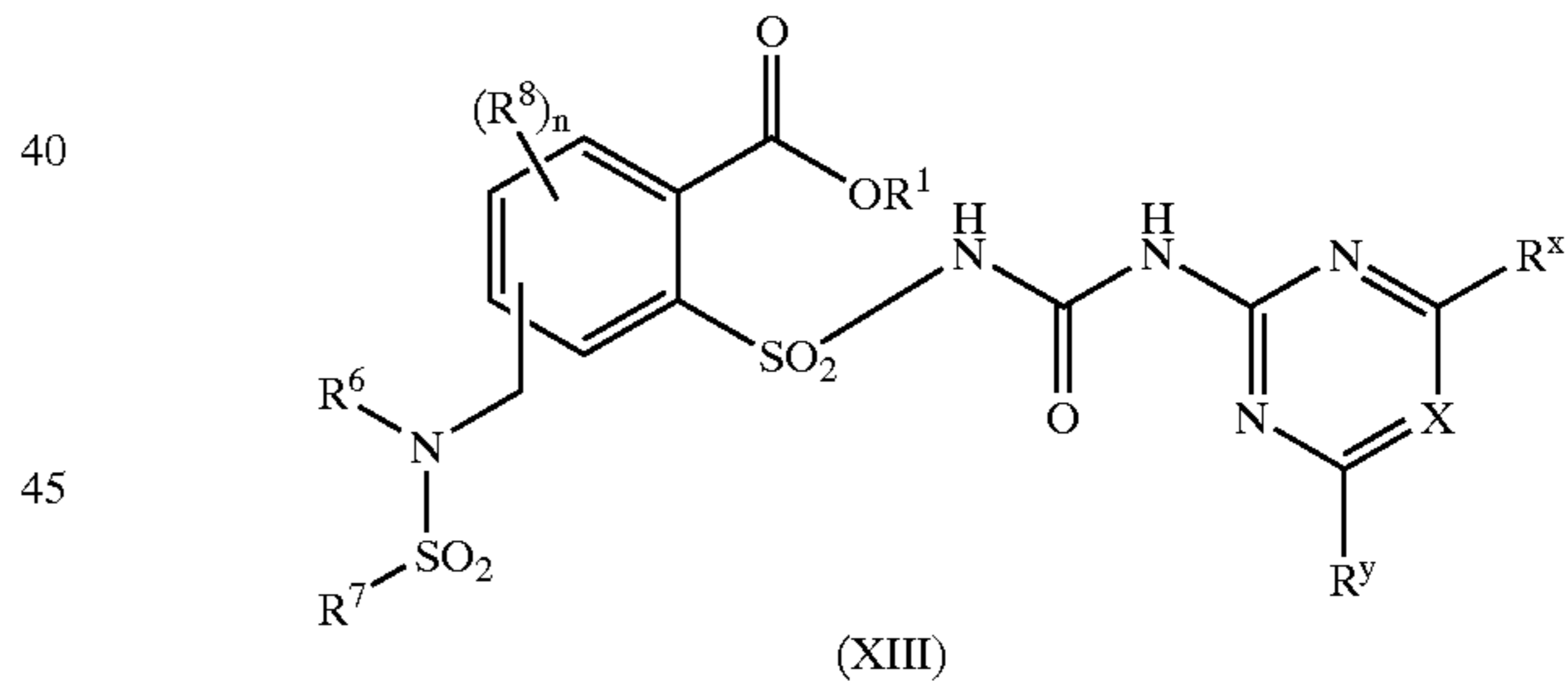
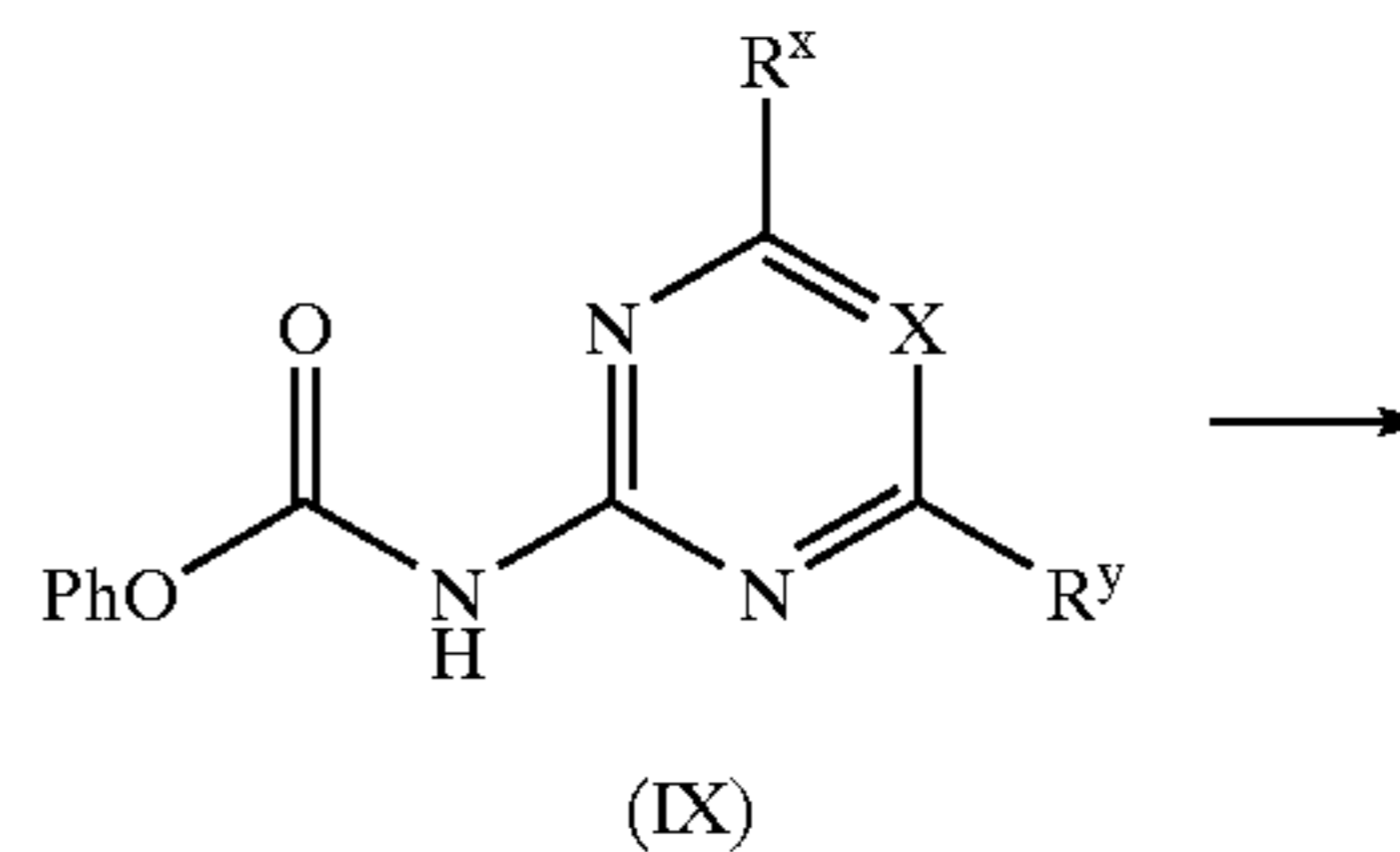
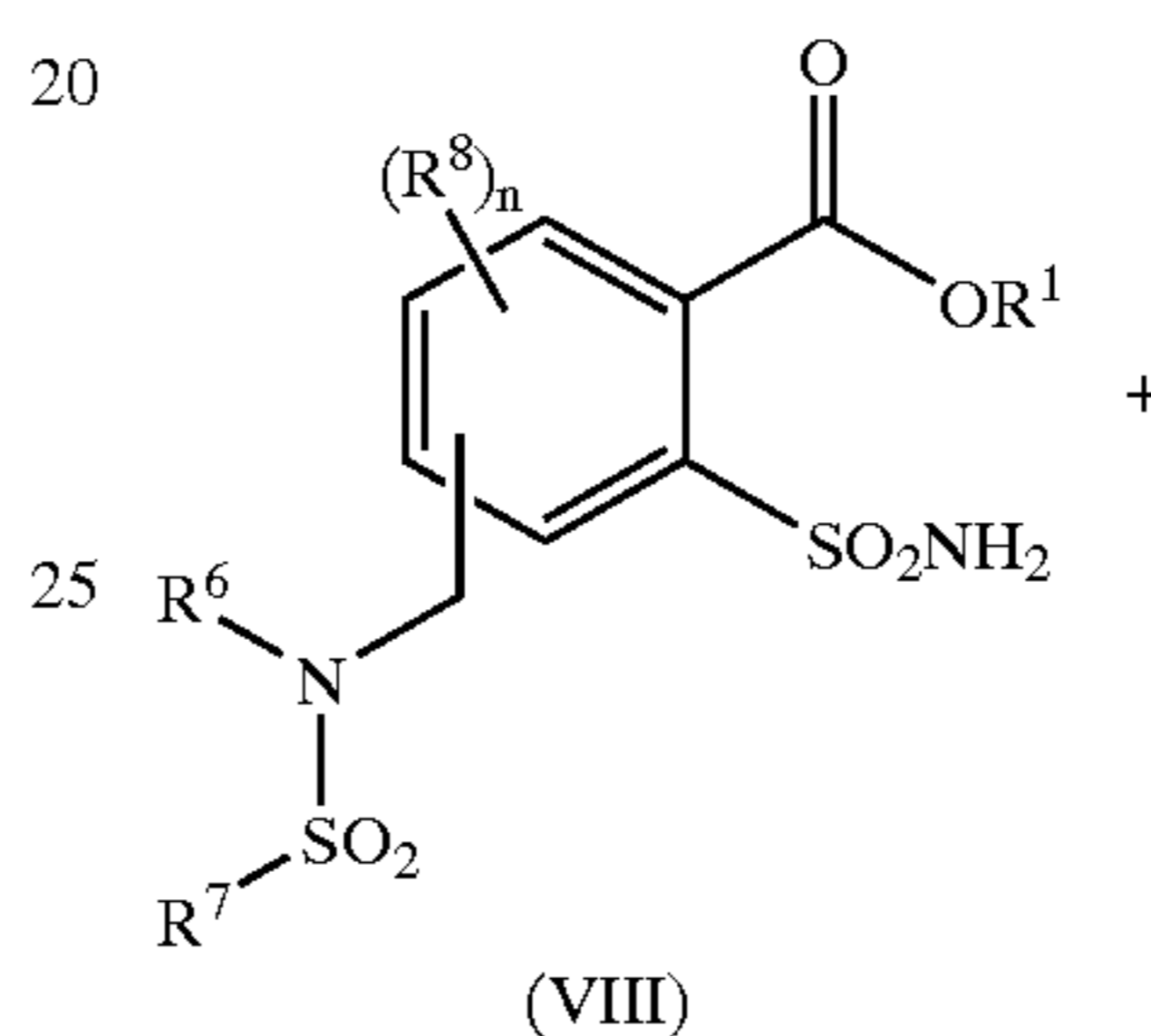


14

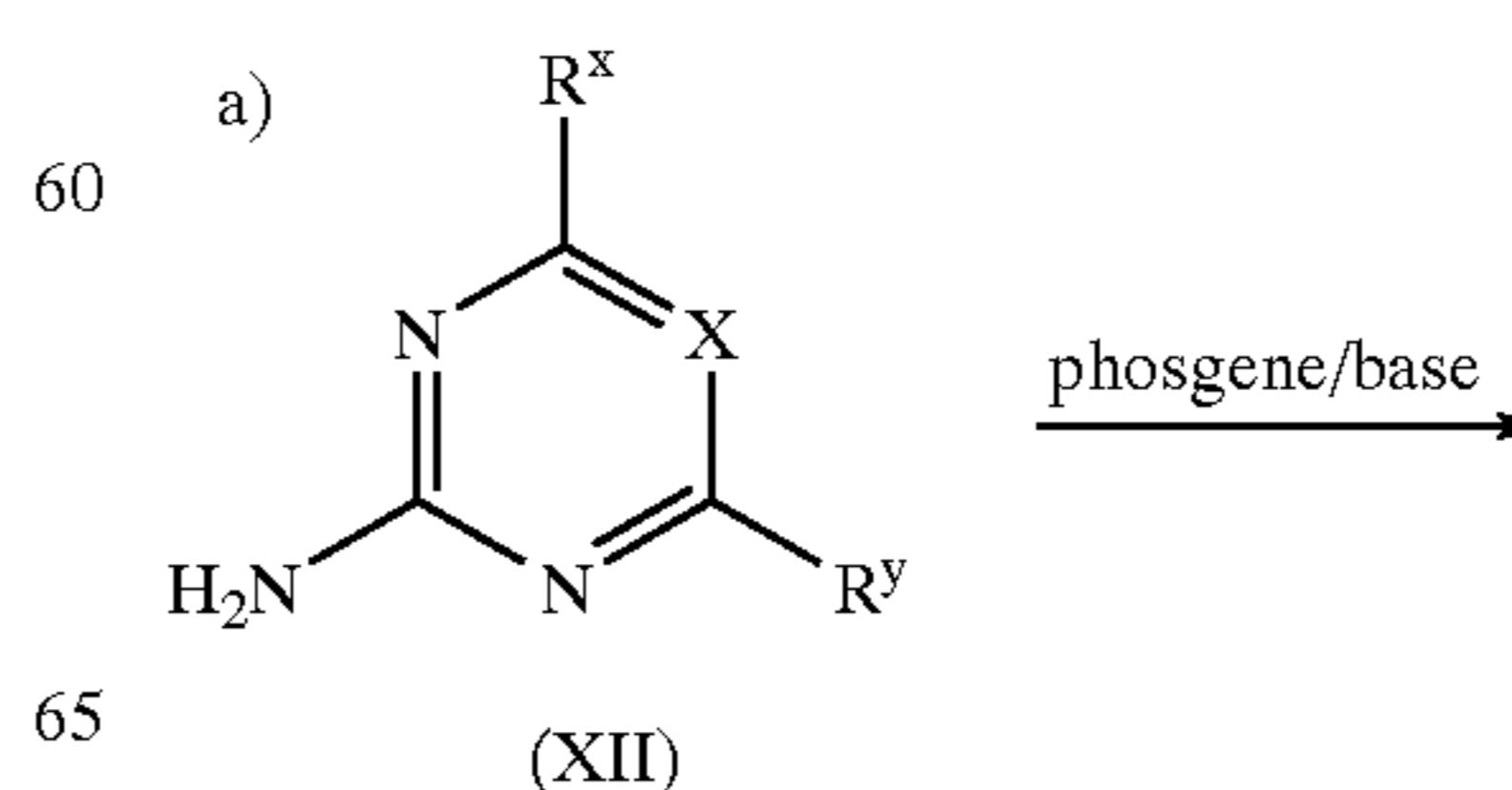
-continued



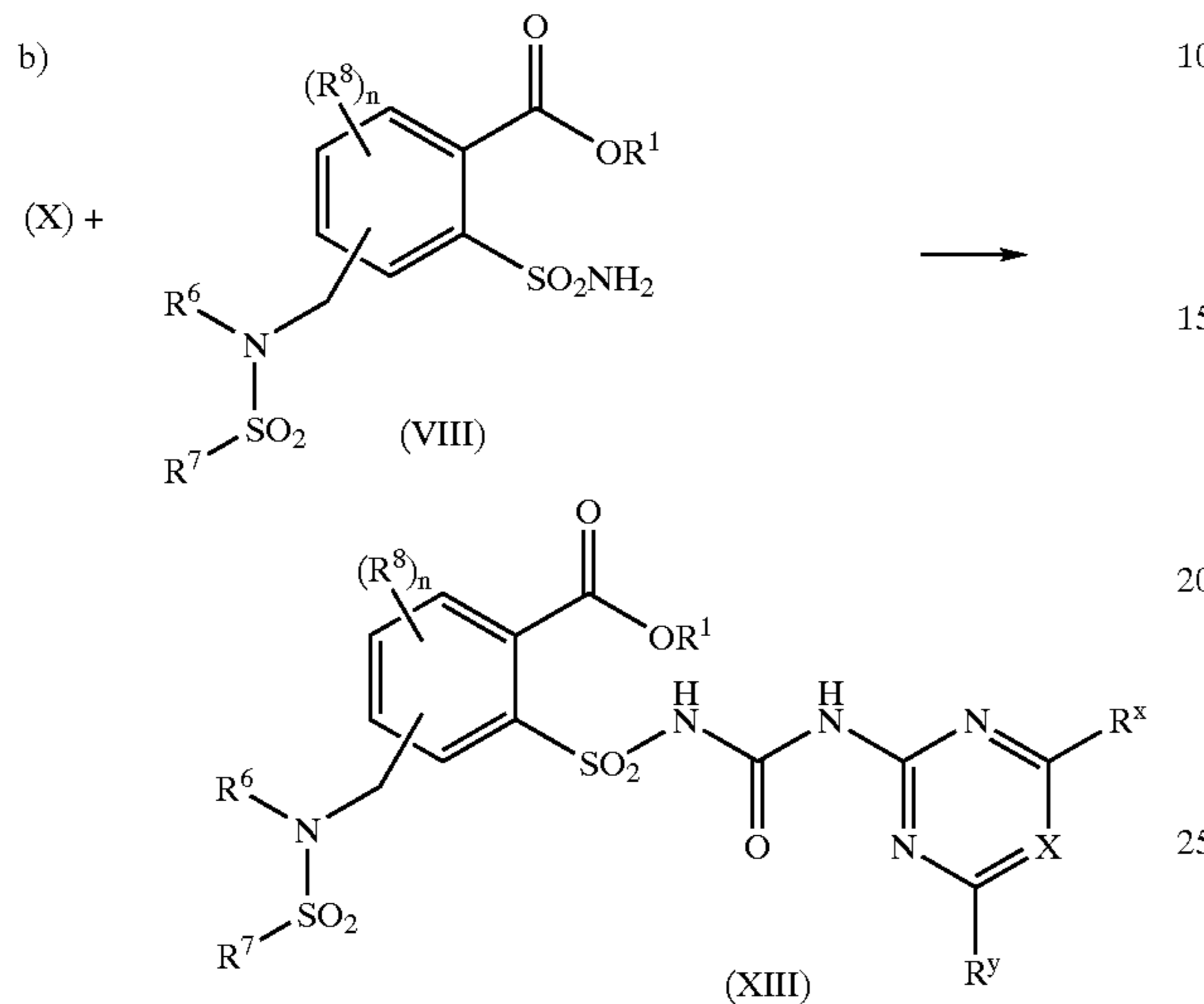
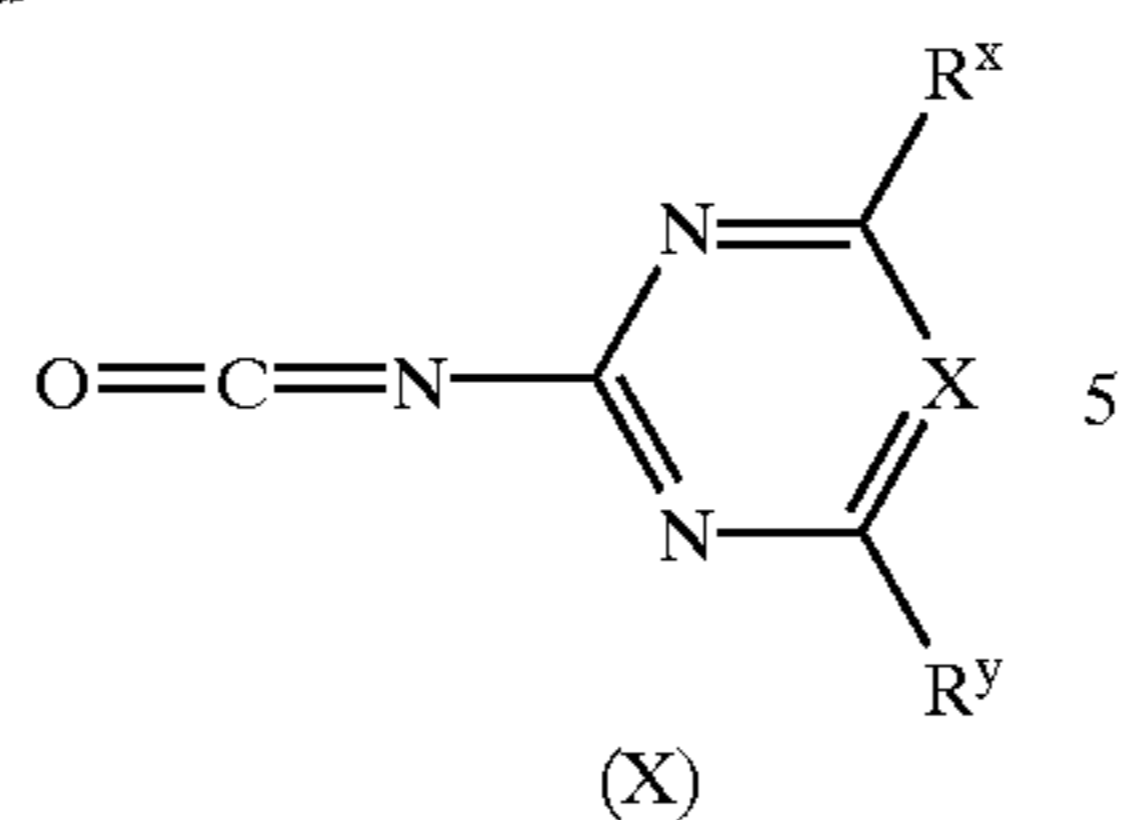
9) reacting a compound of the formula (VIII) with a heterocyclic carbamate of the formula (IX), in which Ph is unsubstituted or substituted phenyl to give the sulfonylurea; or



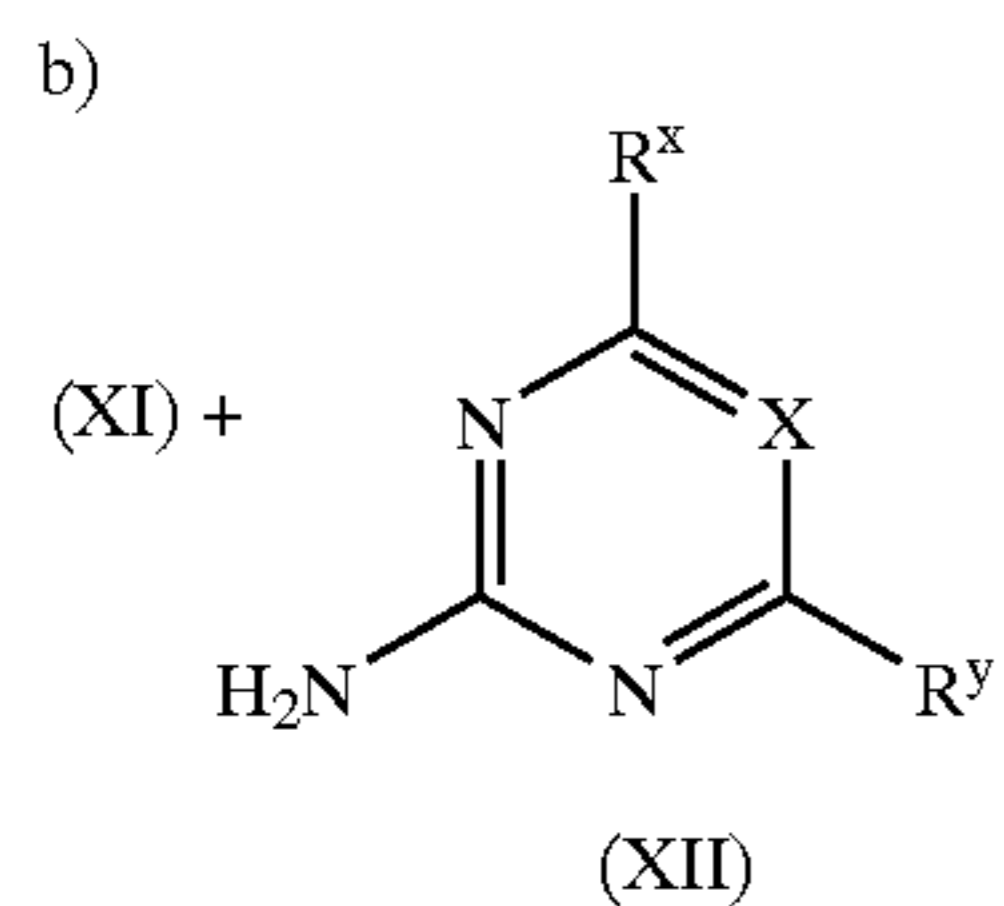
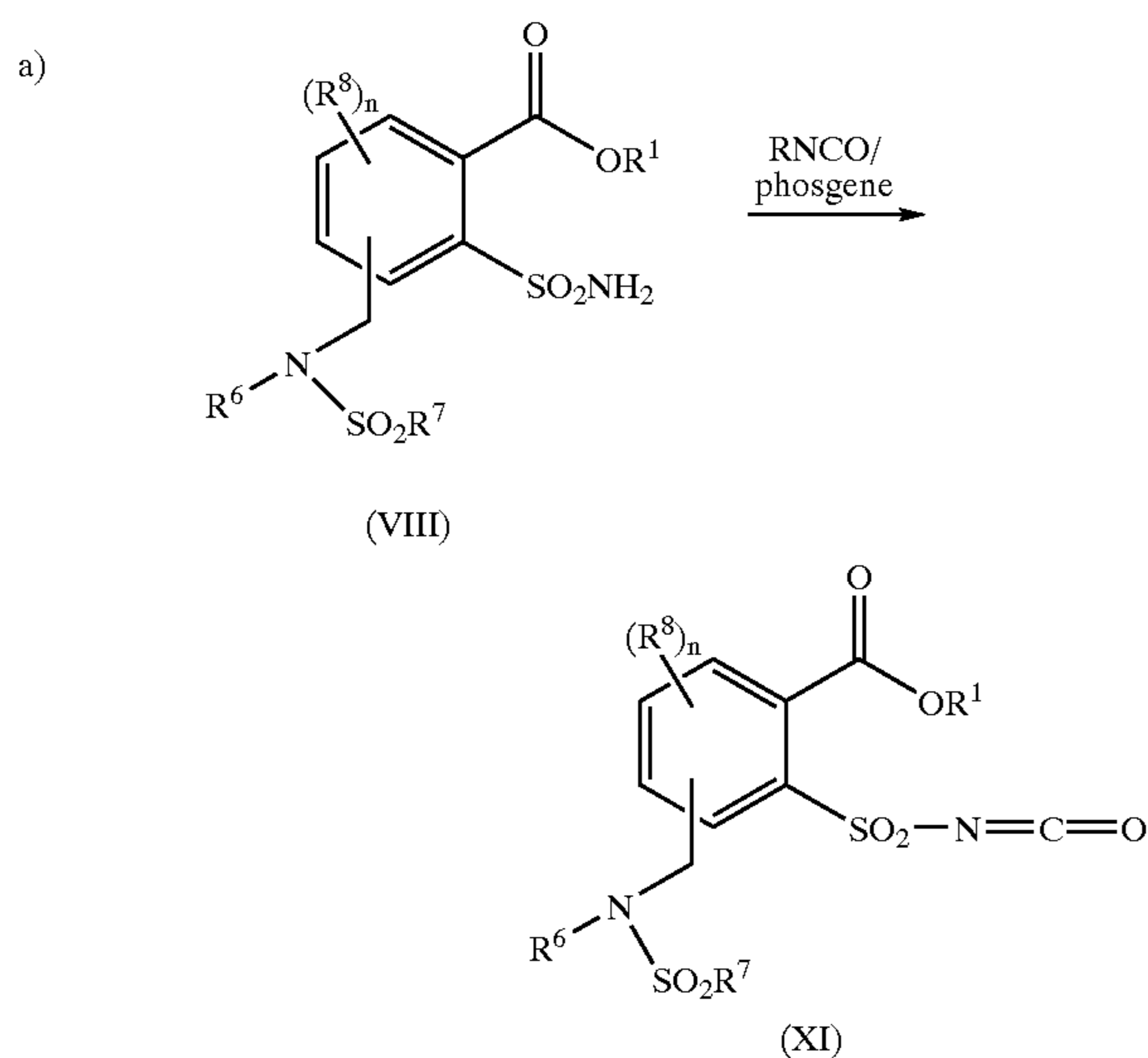
10) a) first reacting an amino heterocycle of the formula (XII) in the presence of a base such as trialkylamine, for example triethylamine, with phosgene to give a heterocyclyl isocyanate of the formula (X), and b) reacting the heterocyclyl isocyanate formed, of the formula (X), with a phenylsulfonamide of the formula (VIII) to give the sulfonylurea; or



15
-continued

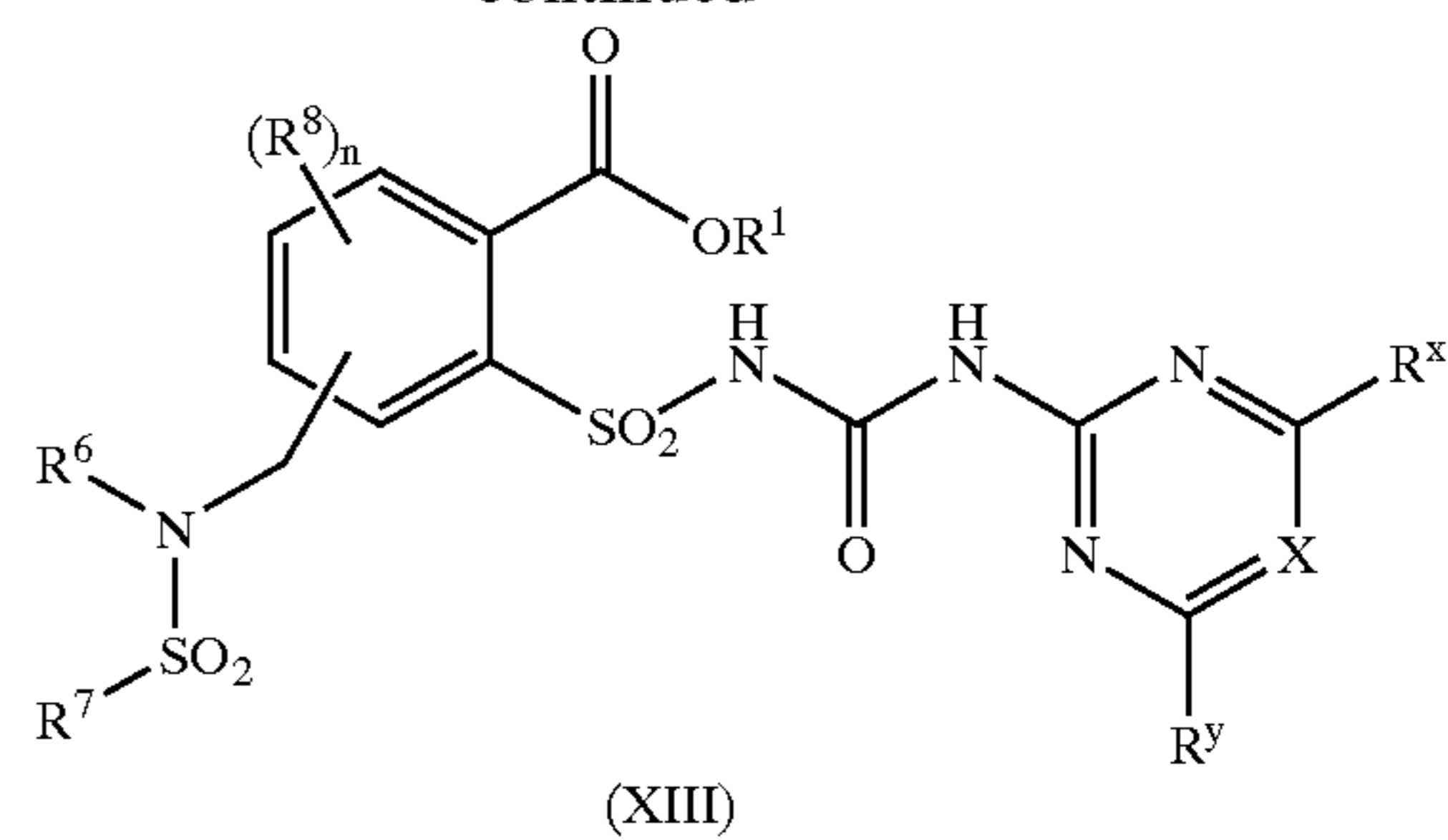


11) a) reacting a compound of the formula (VIII) with an alkyl isocyanate, for example RNCO, in which R=C₁-C₁₀-alkyl and with phosgene to give a sulfonyl isocyanate of the formula (XI), and b) reacting the sulfonyl isocyanate formed, of the formula (XI), with an amino heterocycle of the formula (XII) to give the sulfonamide; or

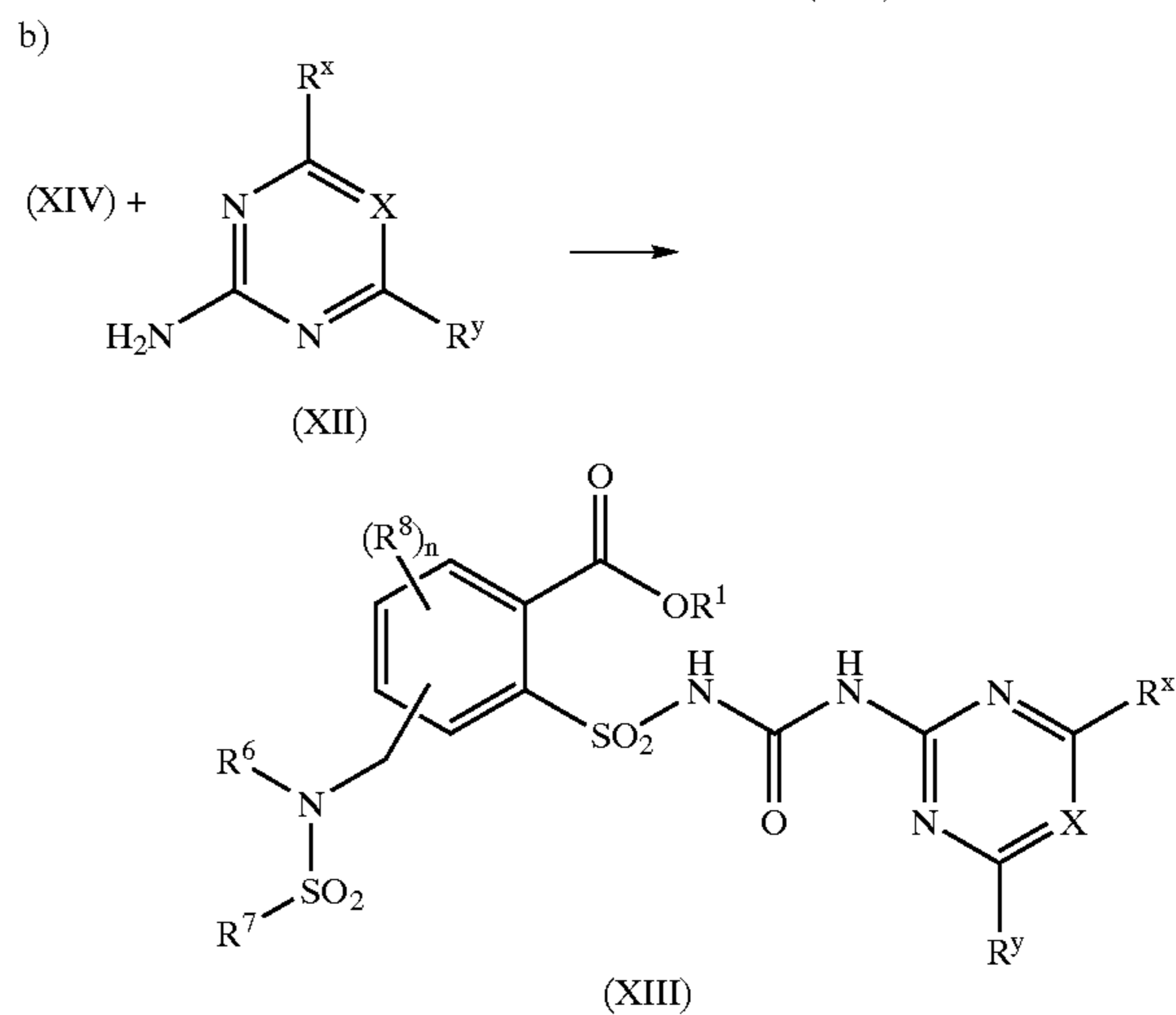
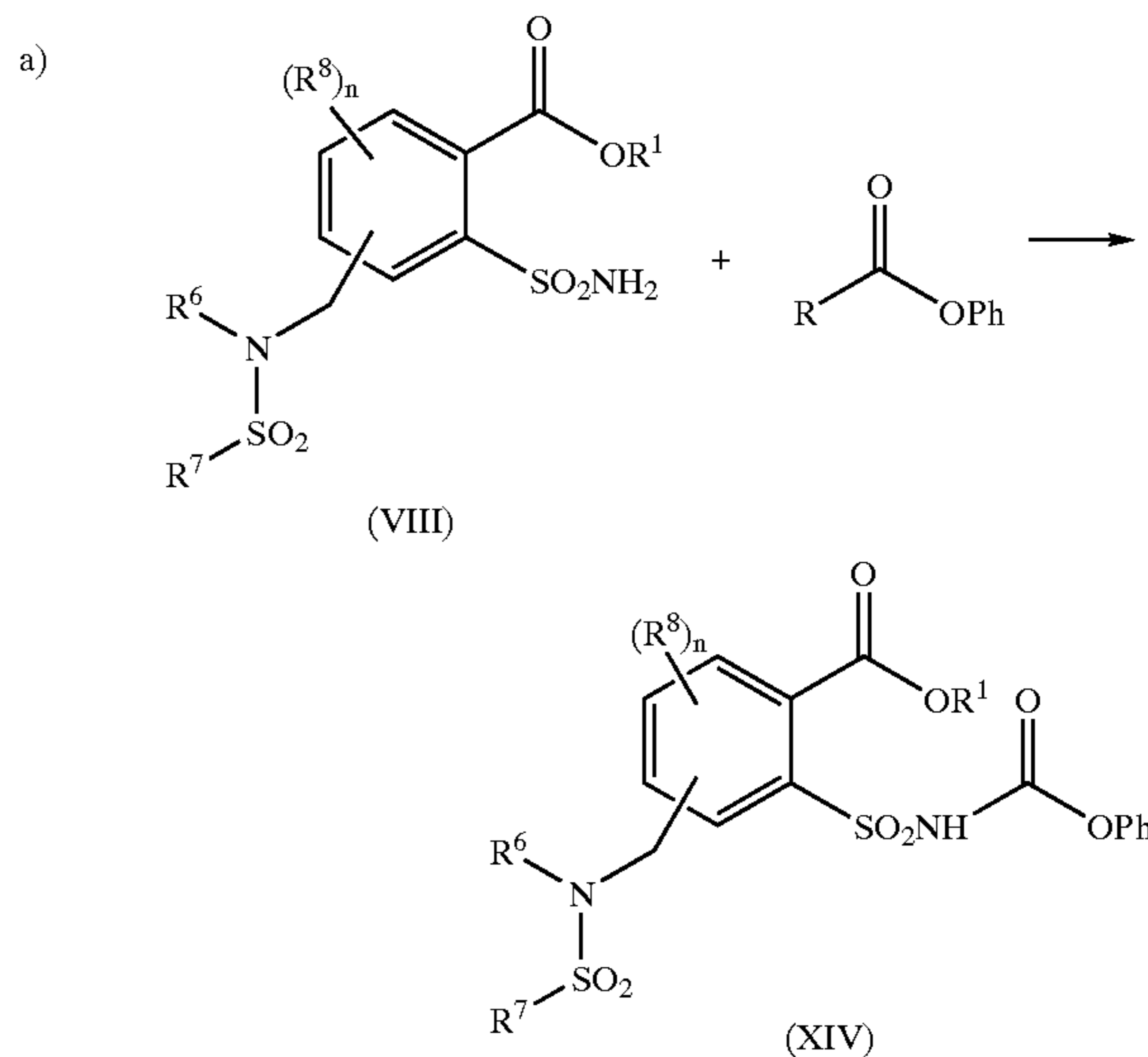


16

-continued



12) a) reacting a compound of the formula (VIII) with a carbonic acid derivative such as R-CO-OPh, in which Ph=unsubstituted or substituted phenyl and R=halogen or unsubstituted or substituted phenoxy to give a phenylsulfonyl carbamate of the formula (XIV), and b) reacting the phenylsulfonyl carbamate formed, of the formula (XIV), in which Ph=unsubstituted or substituted phenyl, with an amino heterocycle of the formula (XII) to give the sulfonamide.



The symbols used in formulae (IX), (X), (XI), (XII), (XIII) and (XIV) have the same meaning as stated in formula (I), including the preferred ranges stated herefor; in addition, the following meanings are also used therein:

R^x , R^y independently of one another are a hydrogen atom, halogen, (C_1-C_4) alkyl, (C_1-C_4) alkoxy, (C_1-C_4) alkylthio, where each of the last-mentioned 3 radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, (C_1-C_4) alkoxy and (C_1-C_4) alkylthio, or are mono- or di $[(C_1-C_4)$ alkyl] amino, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, (C_3-C_6) alkenyloxy or (C_3-C_6) alkynyloxy,

X is CH or N, and

Y is halogen such as fluorine, chlorine, bromine or iodine, preferably chlorine.

Preferred compounds of the formulae (XI), (XIII) and (XIV) are those in which the group $-CH_2-NR^6-SO_2-$ R^7 is in the para position relative to the group $-COOR^1$.

Sulfonylureas such as the compounds of the formula (XIII) can form salts in which the hydrogen of the $-SO_2-NH-$ group is replaced by an agriculturally suitable cation. Examples of these salts are metal salts, in particular alkali metal salts or alkaline earth metal salts, in particular sodium salts and potassium salts, or else ammonium salts or salts with organic amines. Likewise, salt formation can be effected by an addition reaction of an acid with basic groups, such as, for example, amino and alkylamino. Acids which are suitable for this purpose are strong inorganic and organic acids, for example HCl, HBr, H_2SO_4 or HNO_3 . If the present description mentions sulfonylureas such as the compounds of the formula (XIII), this is also to be understood as including their salts in each case.

In process variant 8), the reaction of the sulfonyl halides (VII) is carried out with amino heterocycles of the formula (XII) and cyanates MOCN preferably with base catalysis in inert aprotic organic solvents such as ethyl acetate, tetrahydrofuran, toluene or acetonitrile between $0^\circ C.$ and the boiling point of the solvent. Examples of suitable bases are organic amine bases, in particular pyridines such as pyridine or 3-methylpyridine.

In process variant 9), the reaction of the compounds of the formulae (VIII) and (IX) is carried out preferably with base catalysis in an inert organic solvent such as dichloromethane, acetonitrile, dioxane, tetrahydrofuran or ethyl acetate at between $0^\circ C.$ and the boiling point of the solvent. Examples of bases which are used are K_2CO_3 or organic amine bases such as 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU).

In process variant 10), the reaction of the compound of the formula (XII) is carried out with phosgene to give heterocyclyl isocyanates of the formula (X), for example in inert organic solvents such as ethyl acetate, dioxane or aromatic solvents such as chlorobenzene, if appropriate with addition of an organic amine base such as triethylamine, in general between $0^\circ C.$ and the boiling point of the solvent. The subsequent reaction of the compound of the formula (X) with the compound of the formula (VIII) is carried out for example in inert solvents such as ethyl acetate, dioxane or aromatic solvents such as chlorobenzene, preferably in the presence of bases such as K_2CO_3 or trialkylamines such as triethylamine or tributylamine, in general at temperatures of from $-20^\circ C.$ to the boiling point of the solvent (cf. for example, EP-A-232 067 or EP-A-166516).

In process variant 11), the reaction of the compound of the formula (VIII) is carried out with an alkyl isocyanate and phosgene to give phenylsulfonyl isocyanates of the formula

(XI), for example in inert solvents such as dichloromethane, acetonitrile, dioxane, tetrahydrofuran, toluene or chlorobenzene, in general at temperatures of from $20^\circ C.$ to the boiling point of the solvent. The subsequent reaction of the compound of the formula (XI) with amino heterocycles of the formula (XII) is carried out for example in inert solvents such as dichloromethane, acetonitrile, dioxane, tetrahydrofuran, toluene or chlorobenzene, in general at temperatures of from $0^\circ C.$ to the boiling point of the solvent (cf. for example U.S. Pat. No. 4,481,029).

In process variant 12), the reaction of the compound of the formula (VIII) with a carbonic acid derivative, for example diphenyl carbonate or phenyl chloroformate, to give a phenylsulfonyl carbamate of the formula (XIV) is carried out for example in inert solvents such as xylene, dichloromethane, acetonitrile, dioxane, tetrahydrofuran, toluene or chlorobenzene, preferably in the presence of a base such as K_2CO_3 or organic amine bases such as triethylamine, preferably at temperatures of from $20^\circ C.$ to the boiling point of the solvent (cf., for example, U.S. Pat. No. 4,684,393 and U.S. Pat. No. 4,743,290). The subsequent reaction of the compound of the formula (XIV) with amino heterocycles of the formula (XII) is carried out for example in inert solvents such as xylene, dichloromethane, acetonitrile, dioxane, tetrahydrofuran, toluene or chlorobenzene, in general at temperatures of between $20^\circ C.$ and the boiling point of the solvent.

The compounds of the formula (I) according to the invention thus make possible an efficient preparation of herbicidal sulfonylureas and other active substances.

EXAMPLES

Example 1

a) 3-Amino-4-methoxycarbonylbenzylammonium chloride
After addition of 365 ml of concentrated hydrochloric acid (4.37 mol) and 9 g of PtO_2 , a suspension of 900 g (4.37 mol) of methyl 4-cyano-2-nitrobenzoate in 13.5 l of methanol is first hydrogenated at room temperature at a hydrogen pressure of 1 bar. After the hydrogen uptake has subsided, the pressure is increased to 17 bar, and hydrogenation is continued until the hydrogen uptake is complete. For work-up, the pressure is released to atmospheric pressure, the catalyst is removed by filtration through silica gel and the filtrate is concentrated completely in vacuo. Digestion of the residue with ethyl acetate yields 3-amino-4-methoxycarbonylbenzylammonium chloride, yield 757 g (80%), melting point $185-190^\circ C.$ (decomp.).

b) Methyl 2-amino-4-methanesulfonylaminomethylbenzoate

3 g of 3-amino-4-methoxycarbonylbenzylammonium chloride (18.8 mmol) are dissolved in 50 ml of dimethylacetamide, triethylamine (2.8 g, 27.7 mmol) is added, and the mixture is subsequently reacted at $0-10^\circ C.$ with a solution of methanesulfonyl chloride (1.6 g, 13.8 mmol) in 20 ml of dimethylacetamide. After 1 h, the solvent is removed in vacuo and the residue is worked up by extraction with water/dichloromethane. The combined organic extracts are washed with water and dried (Na_2SO_4) and then evaporated on a rotary evaporator. The residue obtained is crystallized from water, whereupon 3 g (84%) of methyl 2-amino-4-methanesulfonylaminomethylbenzoate of melting point $120-121^\circ C.$ are obtained.

19

c) Methyl 2-chlorosulfonyl-4-methanesulfonylaminomethylbenzoate

After addition of 5 ml of glacial acetic acid, a solution of 3 g (11.6 mmol) of methyl 2-amino-4-methanesulfonylaminomethylbenzoate in 20 ml of concentrated hydrochloric acid is treated at 0–5° C. over 0.5 h with an aqueous NaNO₂ solution (0.81 g, 11.7 mmol, 10 ml of water), and stirring is continued for 0.5 h at 5° C. In parallel, 0.34 g (3.5 mmol) of CuCl is suspended in 30 ml of glacial acetic acid which had previously been saturated with SO₂ gas, and the mixture was subsequently treated with 30 ml of toluene. The diazonium salt solution which had been prepared beforehand is added dropwise to this mixture at 35° C. over 0.5 h, with the evolution of gas starting spontaneously. After 1 hour, the mixture is treated with water, the phases are separated and the aqueous phase is reextracted with dichloromethane. The combined organic phases are washed, dried (Na₂SO₄) and concentrated in vacuo. Extracting the residue by stirring with toluene yields 2.5 g (63%) of methyl 2-chlorosulfonyl-4-methanesulfonylaminomethylbenzoate of melting point 93–94° C.

d) Methyl 2-sulfamoyl-4-methanesulfonylaminomethylbenzoate

A solution of 11 g (32 mmol) of methyl 2-chlorosulfonyl-4-methanesulfonylaminomethylbenzoate in 200 ml of THF is treated at 0° C. with 1.1 g (64 mmol) of NH₃ gas. For work-up, the mixture is concentrated in vacuo. Extracting the residue by stirring with water and then filtration and drying in vacuo give 8.3 g (80%) of methyl 2-sulfamoyl-4-methanesulfonylaminomethylbenzoate of melting point 185–187° C.

e) Methyl 2-[3-(4,6-dimethoxypyrimidin-2-yl)ureidosulfonyl]-4-methanesulfonaminomethylbenzoate

87.15 g (0.2677 mol) of methyl 2-sulfamoyl-4-methanesulfonylaminomethylbenzoate and 74.42 g (0.2677 mol) of N-(4,6-dimethoxypyrimidin-2-yl)phenylcarbamate are suspended in 600 ml of acetonitrile with ice-cooling at 5° C., and the mixture is treated with 40.4 ml (0.2677 mol) of 1,8-diazabicyclo[5.4.0]undec-7-ene in the course of 0.5 hours. After 2 hours at room temperature, approx. 2/3 of the solvent is removed in vacuo and the residue is stirred vigorously with 600 ml of 2N HCl and 400 ml of diisopropyl ether. The product which has precipitated is filtered off with suction, washed in succession with water and diisopropyl ether (in each case twice) and dried in vacuo. This gives 125 g of methyl 2-[3-(4,6-dimethoxypyrimidin-2-yl)ureidosulfonyl]-4-methanesulfonaminomethylbenzoate (92%) of melting point 191–193° C. (decomp.).

Example 2

3-Amino-4-methoxycarbonylbenzylammonium chloride

18.5 g (0.09 mol) of methyl 4-cyano-2-nitrobenzoate and 0.93 g of palladium hydroxide (20% on charcoal) are suspended in a mixture of 8 ml of concentrated hydrochloric acid (30% strength) and 315 ml of water in a Hastelloy stirred autoclave. Then, the mixture is hydrogenated at a hydrogen pressure of 17 bar until the hydrogen uptake is complete. After the pressure in the autoclave has been released and the catalyst filtered off, the filtrate is evaporated completely. This gives 19.5 g (97% strength) of 3-amino-4-methoxycarbonylbenzylammonium chloride of melting point 200–205° C.

20

Example 3

3-Amino-4-methoxycarbonylbenzylammonium chloride

In a stainless steel autoclave, 8.0 g of palladium hydroxide (20% on charcoal) and 100 ml of acetic acid are made inert with N₂. Then, a hydrogen pressure of 17 bar is applied. A solution of 300 g (1.455 mol) of methyl 2-nitro-4-cyanobenzoate in 2.6 l of acetic acid is then metered into the vigorously stirred mixture over 3 hours at +20° C. with cooling, using a metering pump. The H₂ pressure is kept at 17 bar. The pressure in the autoclave is released and the contents are made inert with N₂. The catalyst is filtered off and the filtrates are evaporated. Yield (acetate of the product): 91% of theory in the form of viscous residue. By dissolving the residue in toluene and passing in HCl gas (1 equivalent) at 0 to +10° C., a quantitative precipitate of 3-amino-4-methoxycarbonylbenzylammonium chloride is obtained, which is filtered off in the form of white crystals and dried (yield 91%, melting point 204–206° C.).

Example 4

a) 3-Nitro-4-methoxycarbonylbenzylammonium chloride

700 ml of a 1 M BH₃ solution in THF (0.7 mol) are metered in the course of one hour at 40–50° C. into a solution of 144 g (0.7 mol) of methyl 4-cyano-2-nitrobenzoate in 250 ml of THF and the mixture is subsequently refluxed for a further 1.5 hours. The reaction mixture is then treated with 600 ml of methanol saturated with hydrogen chloride gas, with ice-cooling, and refluxed for one hour. The mixture is concentrated completely under atmospheric pressure and reevaporated with 700 ml of methanol. The residue which remains is stirred with 500 ml of ethyl acetate and filtered, and the product is dried. This gives 115 g (67%) of 3-nitro-4-methoxycarbonylbenzylammonium chloride of melting point 247–248° C.

b) Methyl 4-methanesulfonylaminomethyl-2-nitrobenzoate

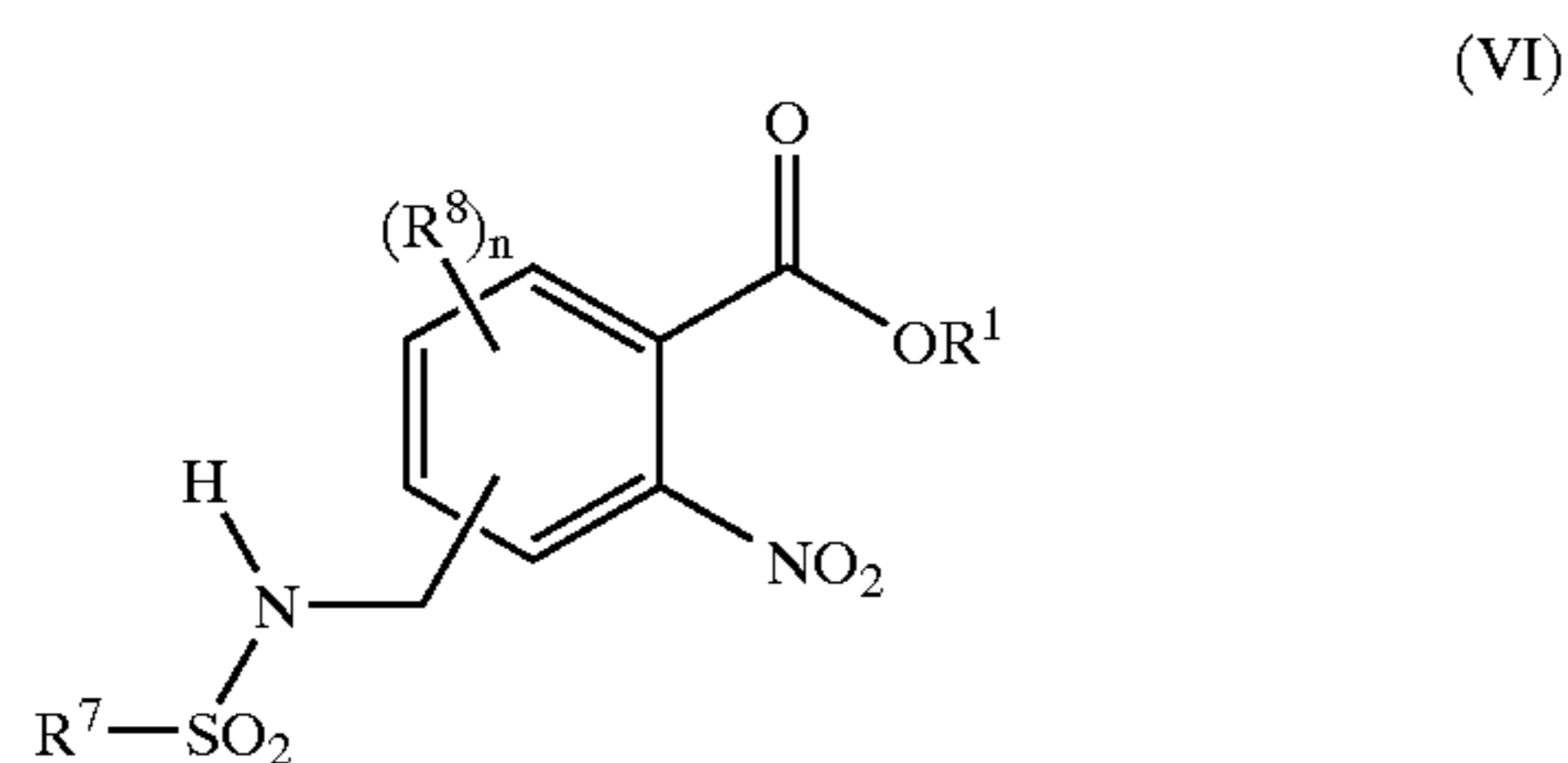
9.4 ml (0.244 mol) of methanesulfonyl chloride are metered over 30 minutes with ice-cooling to a solution of 30.1 g (0.122 mol) of 3-nitro-4-methoxycarbonylbenzylammonium chloride and 34 ml (0.244 mol) of triethylamine in 300 ml of dichloromethane, and stirring of the mixture is continued for one hour. For work-up, the reaction mixture is treated with ice water, the phases are separated, and the aqueous phase is reextracted twice more with dichloromethane. After drying (Na₂SO₄), filtration and concentration, 31.3 g (89%) of methyl 4-methanesulfonylaminomethyl-2-nitrobenzoate remain as a syrupy oil.

c) Methyl 2-amino-4-methanesulfonylaminomethylbenzoate 13.25 g (46 mmol) of methyl 4-methanesulfonylaminomethyl-2-nitrobenzoate are dissolved in 200 ml of methanol. After addition of 1.4 ml (46 mmol) of concentrated hydrochloric acid and 1 g of Pd catalyst (10% on active charcoal), the mixture is hydrogenated with hydrogen under atmospheric pressure until the hydrogen uptake is complete. The mixture is removed from the catalyst by filtration and the filtrate is concentrated completely. Crystallization of the residue from water yields 10.4 g (88%) of methyl 2-amino-4-methanesulfonylaminomethylbenzoate of melting point 121–122° C.

21

We claim:

1. A compound of the formula



in which

R^1 is H, (C_1-C_8) alkyl, (C_3-C_8) alkenyl or (C_3-C_8) alkynyl, where the last 3 radicals are unsubstituted or substituted,

R^6 is H or (C_1-C_8) alkyl which is unsubstituted or substituted,

R^7 is (C_1-C_8) alkyl, (C_3-C_8) alkenyl, (C_3-C_8) alkynyl, (C_6-C_{14}) aryl or mono- or di- (C_1-C_8) alkylamino which are unsubstituted or substituted, or

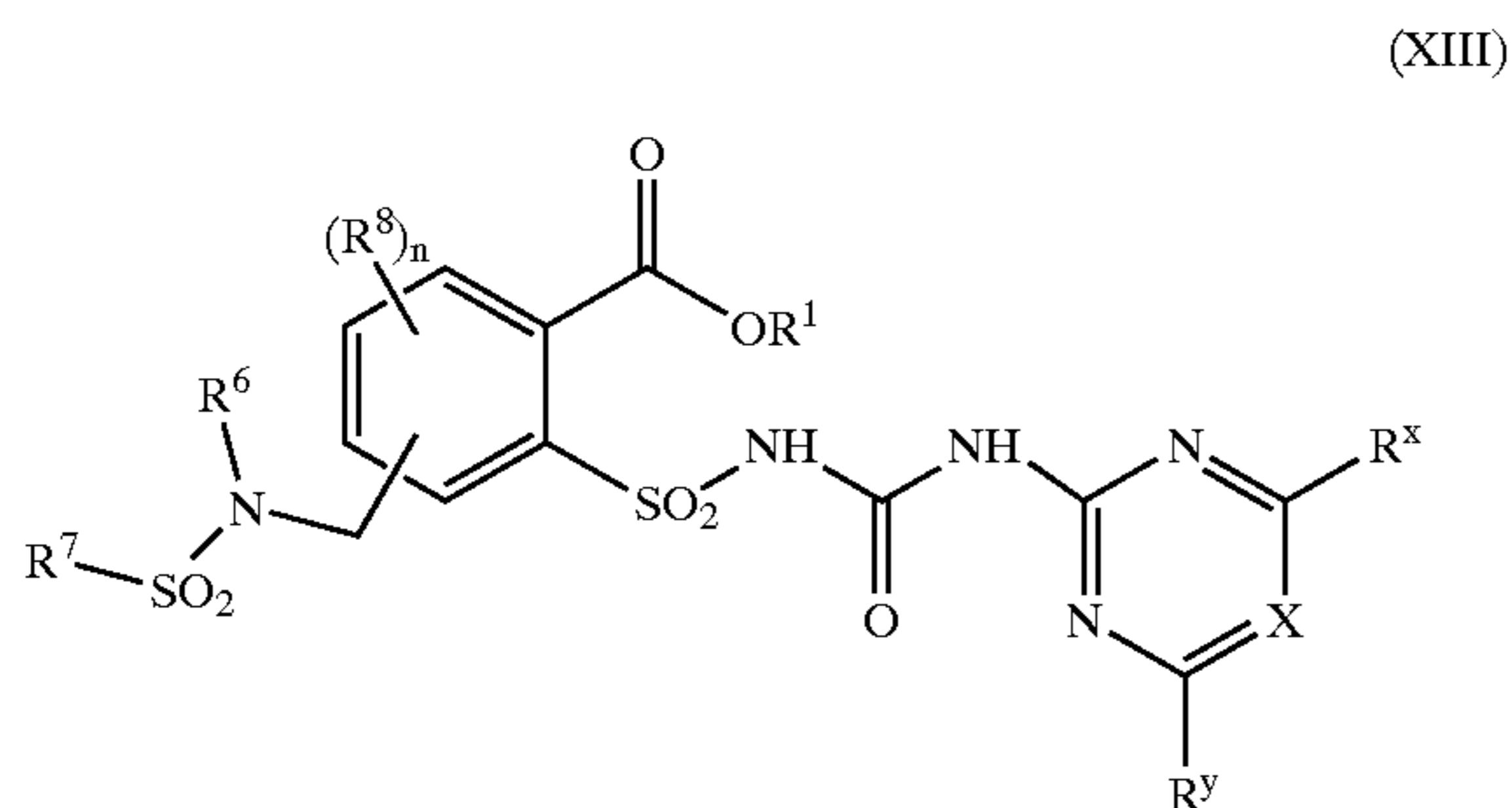
R^6 and R^7 together form a chain of the formula $-(CH_2)_m E_m^1-$, which is unsubstituted or substituted, and where $m=2, 3$ or 4 , $m^1=0$ or 1 and $E=CO$ or SO_2 ,

R^8 radicals are identical or different and are (C_1-C_4) alkyl, (C_1-C_4) alkoxy, $[(C_1-C_4)$ alkyl]carbonyl or $[(C_1-C_4)$ alkoxy]carbonyl which are unsubstituted or substituted, or R^8 is halogen, NO_2 or CN ,

n is 0, 1, 2 or 3, and

wherein, the substituents are halogen, (C_1-C_8) alkoxy, (C_1-C_8) haloalkoxy, (C_1-C_8) alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido, (C_1-C_8) alkoxycarbonyl, (C_1-C_8) alkylcarbonyl, formyl, carbamoyl, mono- and di- (C_1-C_8) alkylaminocarbonyl, acylamino, mono- and di- (C_1-C_8) alkylamino, (C_1-C_8) alkylsulfinyl, (C_1-C_8) haloalkylsulfinyl, (C_1-C_8) alkylsulfonyl, halo- (C_1-C_8) alkylsulfonyl, (C_1-C_8) cycloalkyl, (C_1-C_8) cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C_1-C_8) alkyl and halo- (C_1-C_8) alkyl, (C_1-C_8) alkenyl, (C_1-C_8) alkynyl, (C_1-C_8) alkenyloxy or (C_1-C_8) alkynyloxy.

2. A process for preparing a phenylsulfonylurea of the formula



in which

R^1 is H, (C_1-C_8) alkyl, (C_3-C_8) alkenyl or (C_3-C_8) alkynyl, where the last 3 radicals are unsubstituted or substituted,

R^6 is H or (C_1-C_8) alkyl which is unsubstituted or substituted,

R^7 is (C_1-C_8) alkyl, (C_3-C_8) alkenyl, (C_3-C_8) alkynyl, (C_6-C_{14}) aryl or mono- or di- (C_1-C_8) alkylamino which are unsubstituted or substituted, or

22

R^6 and R^7 together form a chain of the formula $-(CH_2)_m E_m^1-$ which is unsubstituted or substituted, and where $m=2, 3$ or 4 , $m^1=0$ or 1 and $E=CO$ or SO_2 ,

R^8 radicals are identical or different and are (C_1-C_4) alkyl, (C_1-C_4) alkoxy, $[(C_1-C_4)$ alkyl]carbonyl or $[(C_1-C_4)$ alkoxy]carbonyl which are unsubstituted or substituted, or R^8 is halogen, NO_2 or CN , and

n is 0, 1, 2 or 3,

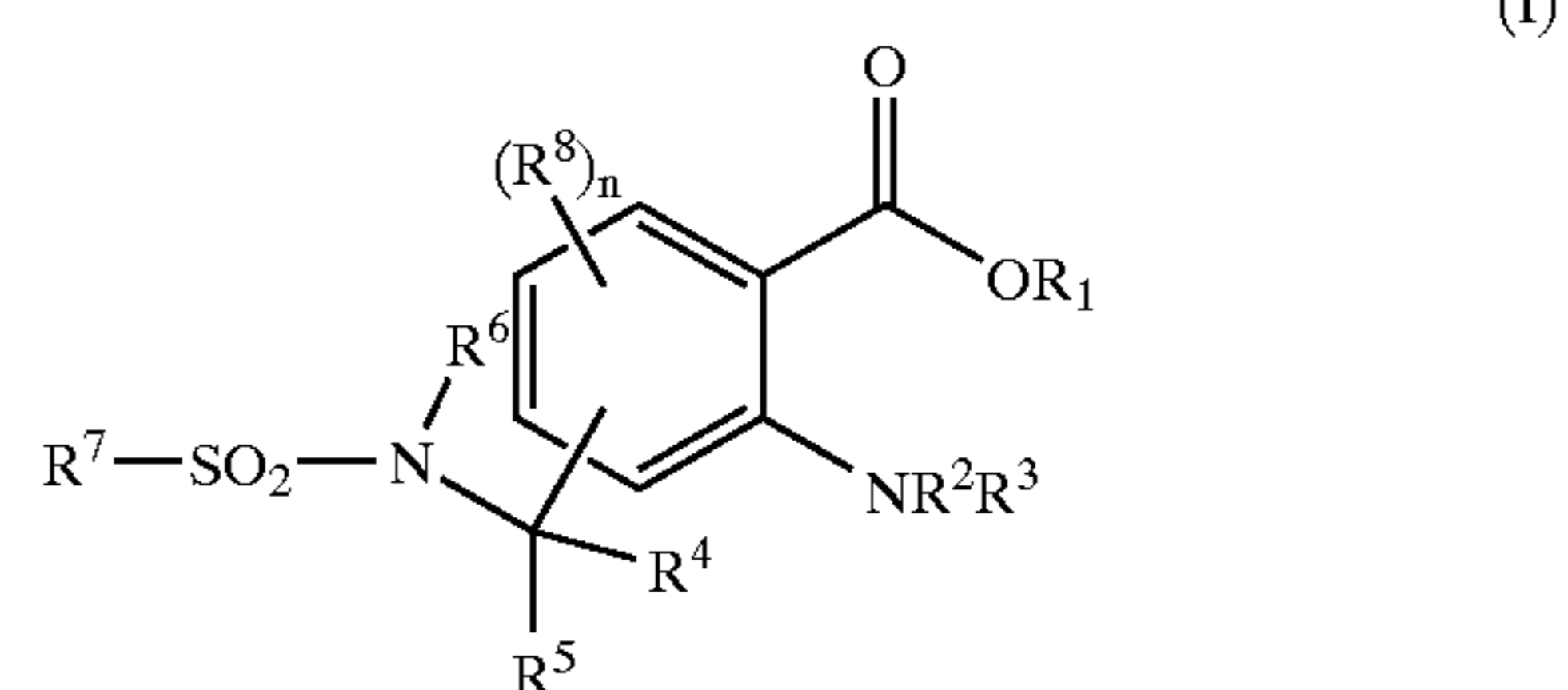
R^x, R^y independently of one another are a hydrogen atom, halogen, (C_1-C_4) alkyl, (C_1-C_4) alkoxy, (C_1-C_4) alkylthio, where each of the last-mentioned 3 radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, (C_1-C_4) alkoxy and (C_1-C_4) alkylthio, or are mono- or di- $[(C_1-C_4)$ alkyl]amino, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, (C_3-C_6) alkenyloxy or (C_3-C_6) alkynyloxy, and

X is CH or N

and wherein, unless otherwise identified, the substituents are halogen, (C_1-C_8) alkoxy, (C_1-C_8) haloalkoxy, (C_1-C_8) alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido, (C_1-C_8) alkoxycarbonyl, (C_1-C_8) alkylcarbonyl, formyl, carbamoyl, mono- and di- (C_1-C_8) alkylaminocarbonyl, acylamino, mono- and di- (C_1-C_8) alkylamino, (C_1-C_8) alkylsulfinyl, (C_1-C_8) haloalkylsulfinyl, (C_1-C_8) alkylsulfonyl, halo- (C_1-C_8) alkylsulfonyl, (C_1-C_8) cycloalkyl, (C_1-C_8) cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C_1-C_8) alkyl and halo- (C_1-C_8) alkyl, (C_1-C_8) alkenyl, (C_1-C_8) alkynyl, (C_1-C_8) alkenyloxy or (C_1-C_8) alkynyloxy;

which comprises:

a) reacting a compound of formula (VI) according to claim 1 with a reagent that reduces a nitro group to an amine or by catalytic hydrogenation to give a compound of the formula

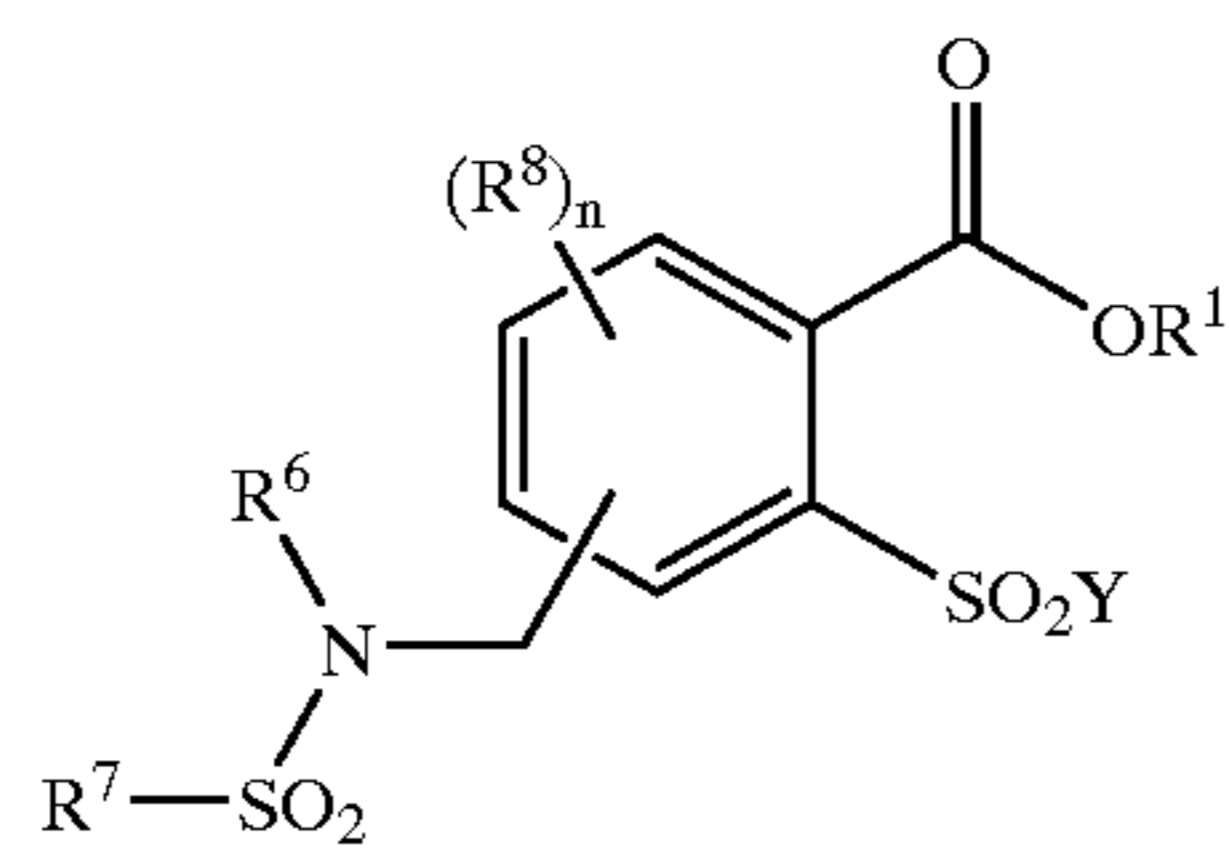


which R^2, R^3 , and R^6 are hydrogen

b) reacting the compound of formula (I) with an alkylating agent or by reductive amination to give a compound of the formula (I) where R^6 is an unsubstituted or substituted C_1-C_8 -alkyl group

c) first reacting the compound formed in step b) with a diazotizing agent in the presence of an acid followed by an SO_2 source in the presence of a copper catalyst and an acid to give a compound of the formula

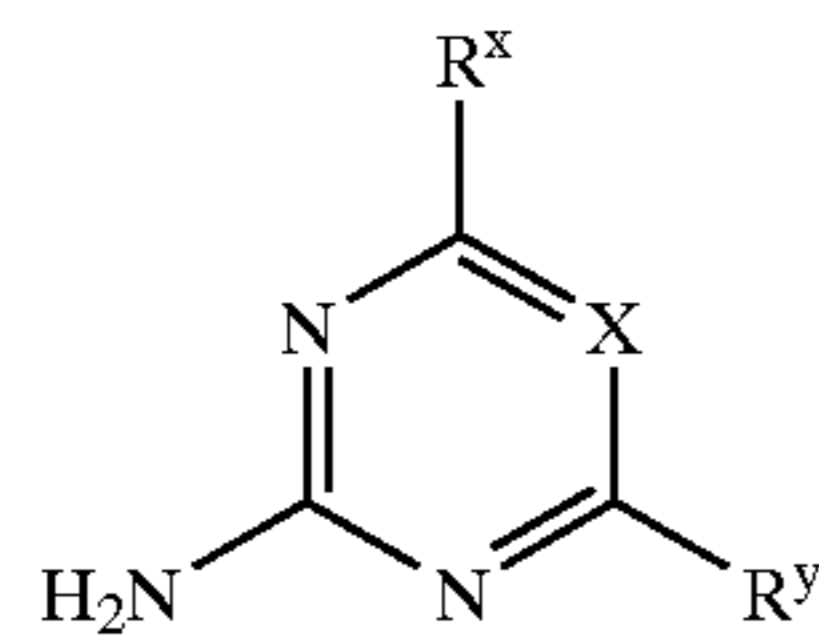
23



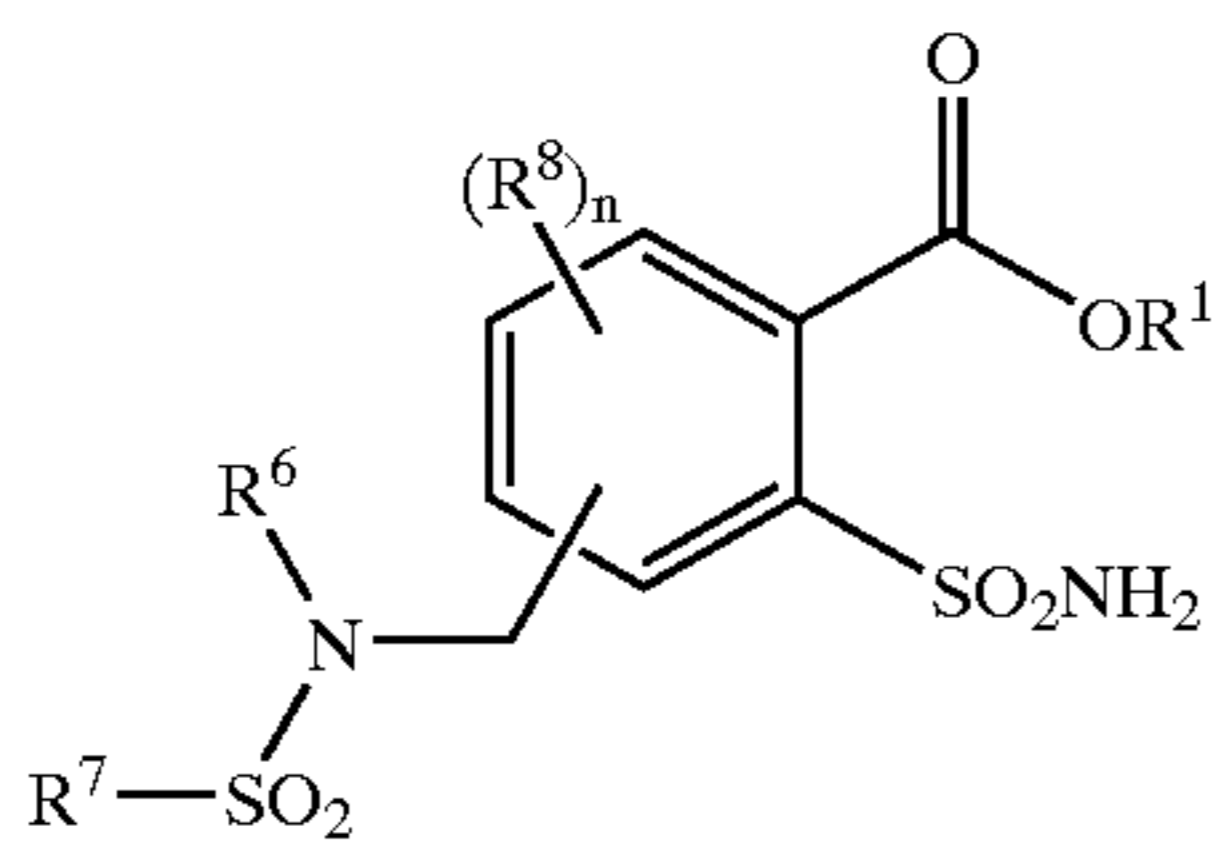
where Y is halogen

d) reacting the compound of formula (VII) with either:

i) amine of the formula (XII) in the presence of MOCN, wherein M is an ammonium ion or an alkali metal ion, to give a compound of the formula (XIII)

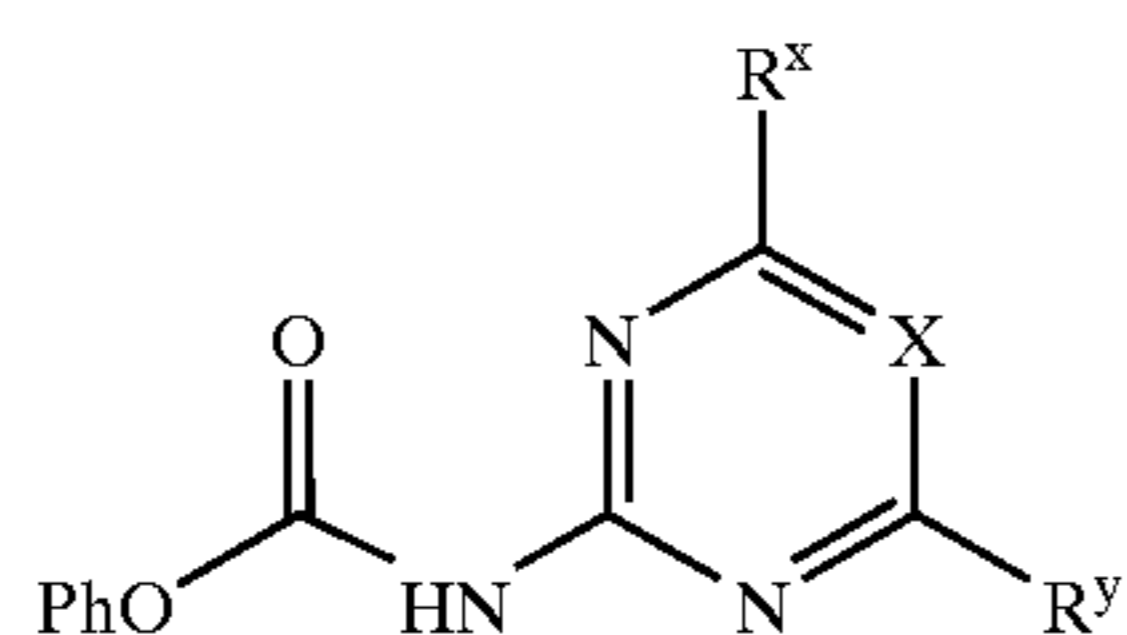


ii) ammonia in the presence of a suitable solvent to give a compound



and subsequently preparing the compound of the formula (XIII) by:

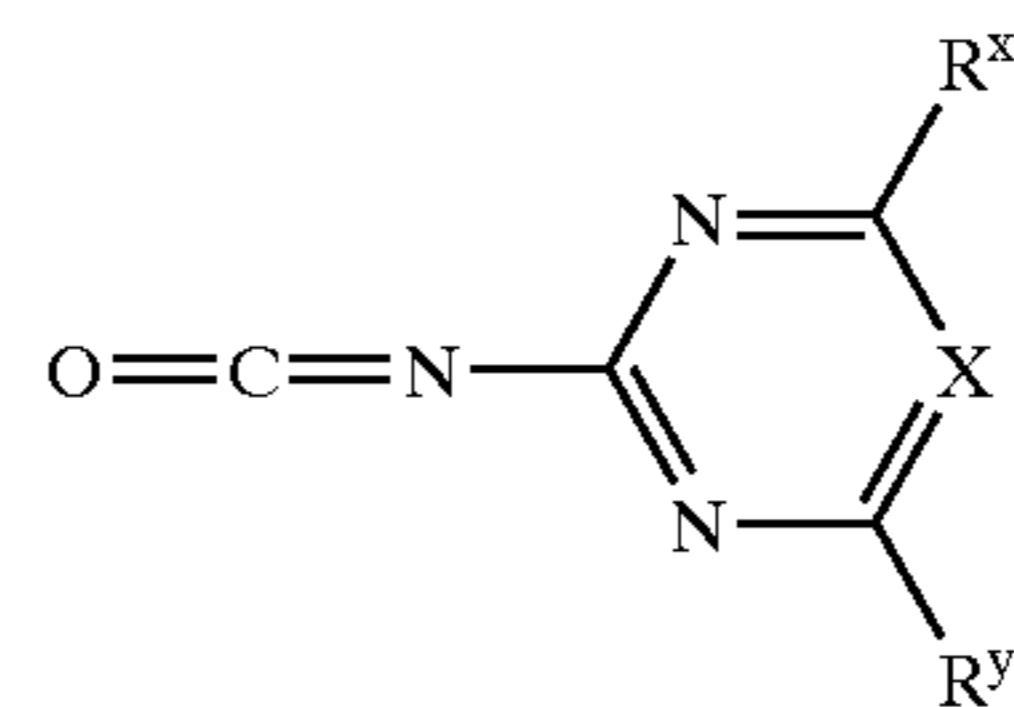
a) reacting the compound of the formula (VIII) with a compound of the formula (IX) to give the compound of the formula (XII)



or

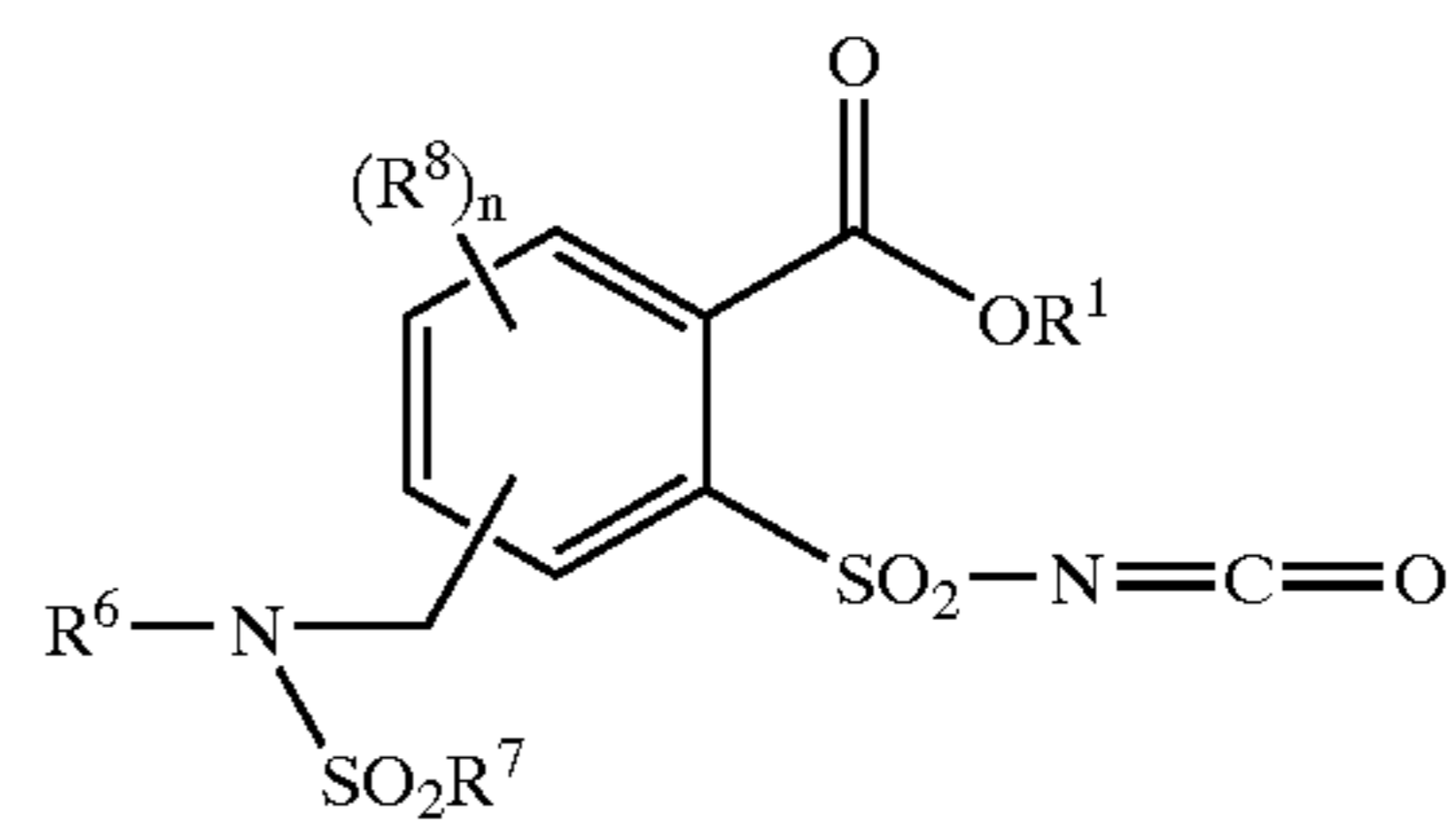
b) reacting the compound of the formula (VIII) with an isocyanate of the formula (X) to give the compound of the formula (XIII)

24

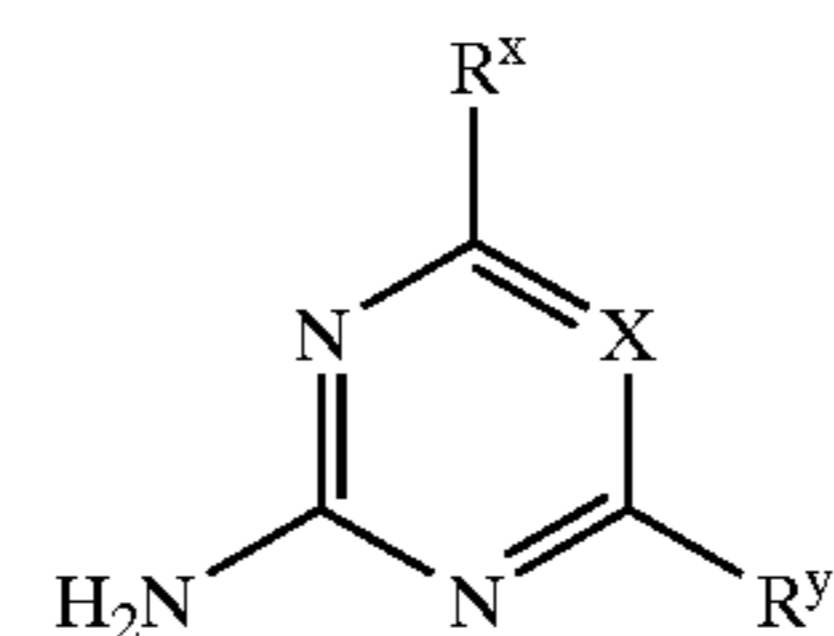


or

c) reacting the compound of the formula (VIII) with an alkyl isocyanate and phosgene to give a compound of the formula (XI)

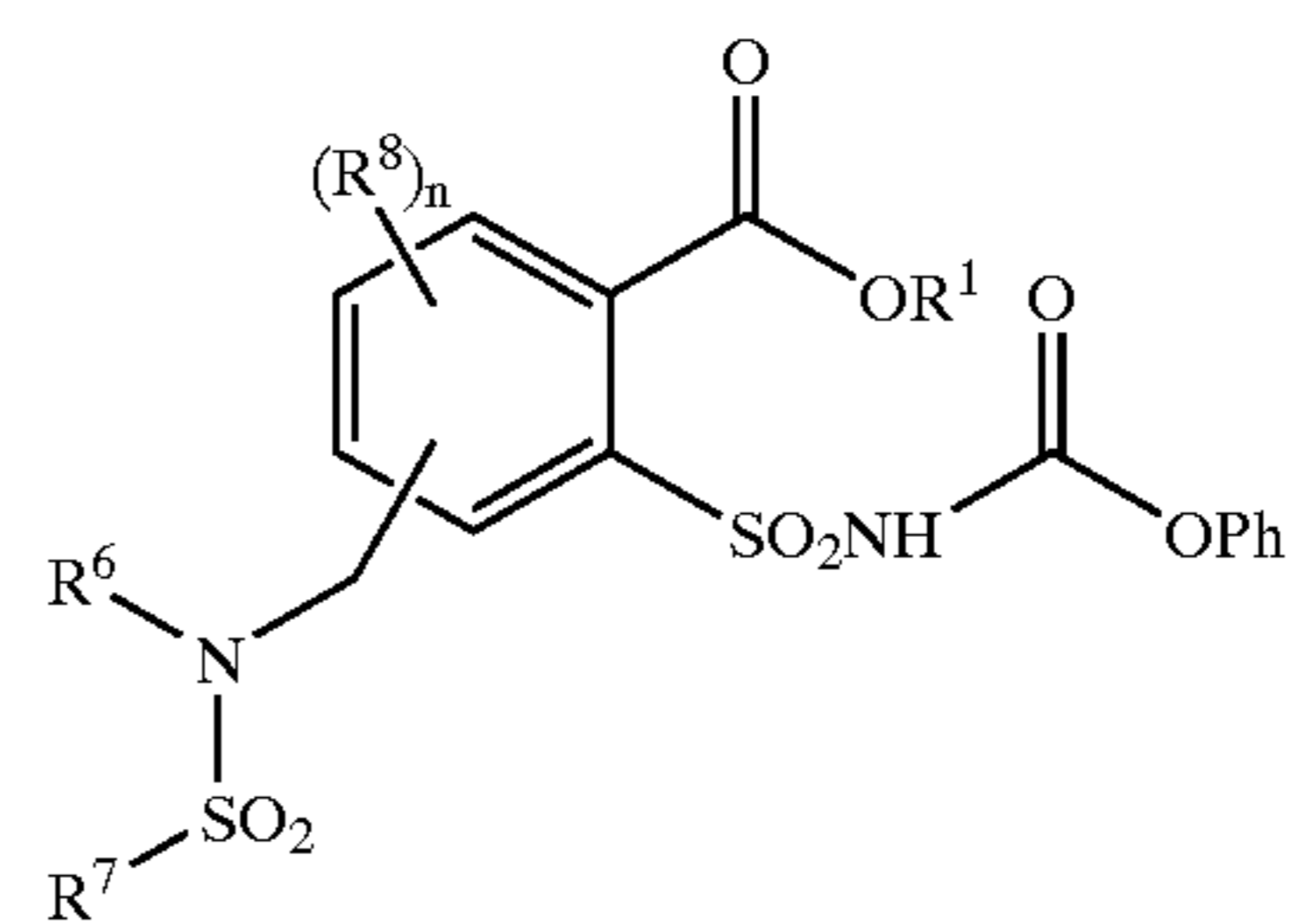


which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

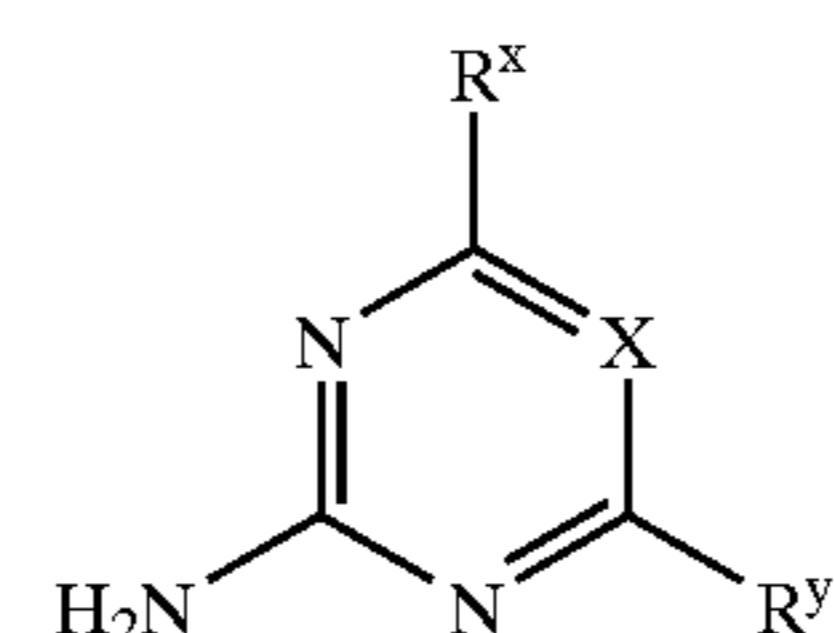


or

d) reacting the compound of the formula (VIII) with a carbonic acid derivative R-CO-OPh, in which Ph=unsubstituted or substituted phenyl and R=halogen or unsubstituted or substituted phenoxy to give a compound of the formula (XIV)



which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

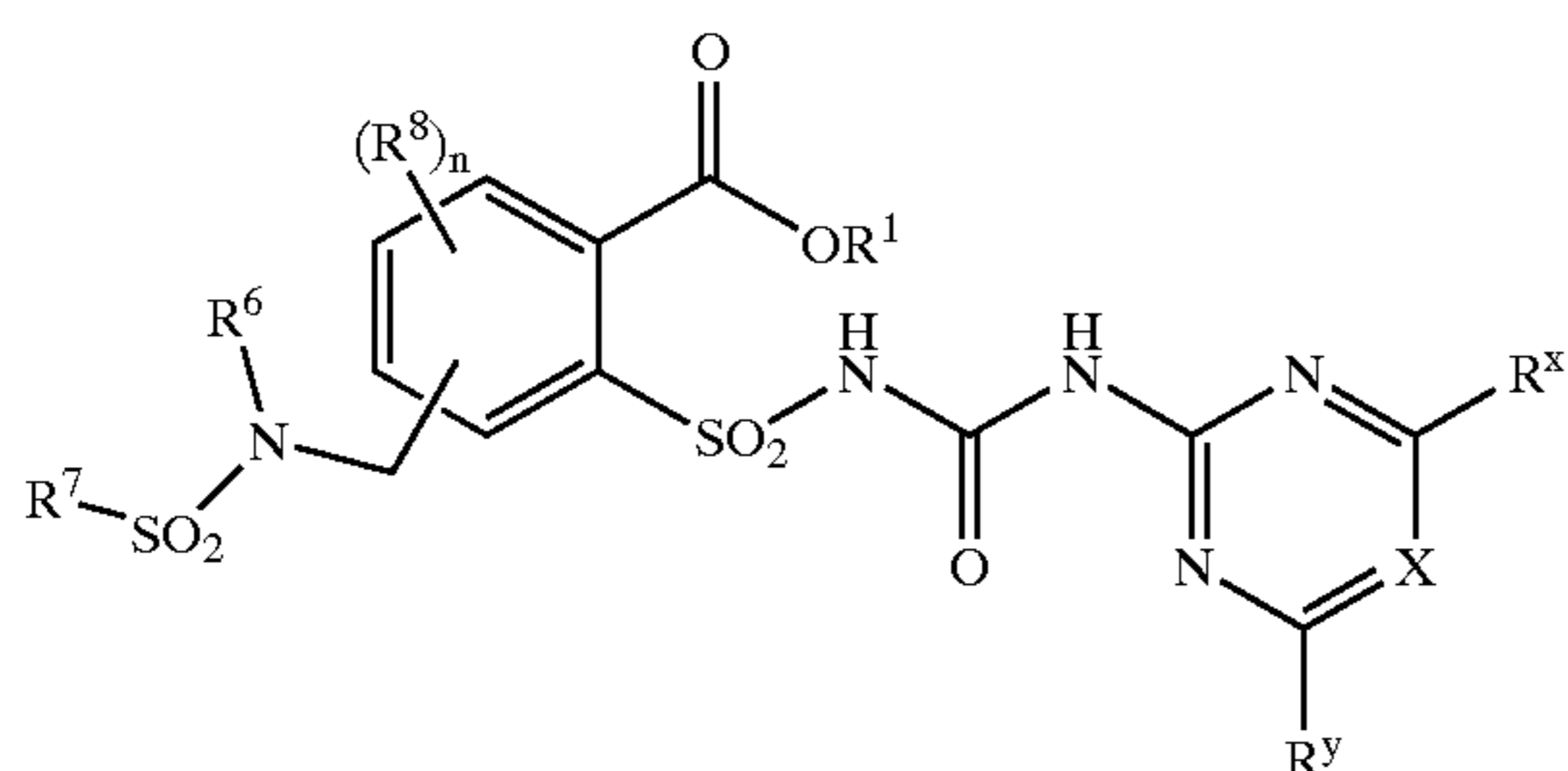


where R¹, R⁷, R⁸, and n in formulae (XI) and (XIV) are as defined in formula (I) R^x, R^y and X in formulae (IX), (X) and

25

(XII) are as defined in formula (XIII) of the present claim and Ph in formulae (IX) and (XIV) is unsubstituted or substituted phenyl, wherein the substituents on the phenyl ring are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₁-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C₁-C₈)alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy.

3. A process of preparing a sulfonylurea of the formula



(XIII)

in which

R¹ is H, (C₁-C₈)alkyl, (C₃-C₈)alkenyl or (C₃-C₈)alkynyl, where the last 3 radicals are unsubstituted or substituted,

R⁶ is H (C₁-C₈)alkyl which is unsubstituted or substituted,

R⁷ is (C₁-C₈)alkyl, (C₃-C₈)alkenyl, (C₃-C₈)alkynyl, (C₆-C₁₄)aryl or mono- or di(C₁-C₈)alkylamino which are unsubstituted or substituted, or

R⁶ and R⁷ together form a chain of the formula —(CH₂)_mE_m¹— which is unsubstituted or substituted, and where m=2, 3 or 4, m¹=0 or 1 and E=CO or SO₂,

R⁸ radicals are identical or different and are (C₁-C₄)alkyl, (C₁-C₄)alkoxy, [(C₁-C₄)alkyl]carbonyl or [(C₁-C₄)alkoxy]carbonyl which are unsubstituted or substituted, or R⁸ is halogen, NO₂ or CN, and

n is 0, 1, 2 or 3,

R^x, R^y independently of one another are a hydrogen atom, halogen, (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)alkylthio, where each of the last-mentioned 3 radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, (C₁-C₄)alkoxy and (C₁-C₄)alkylthio, or are mono- or di[(C₁-C₄)alkyl]amino, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, (C₃-C₆)alkenyloxy or (C₃-C₆)alkynyloxy, and

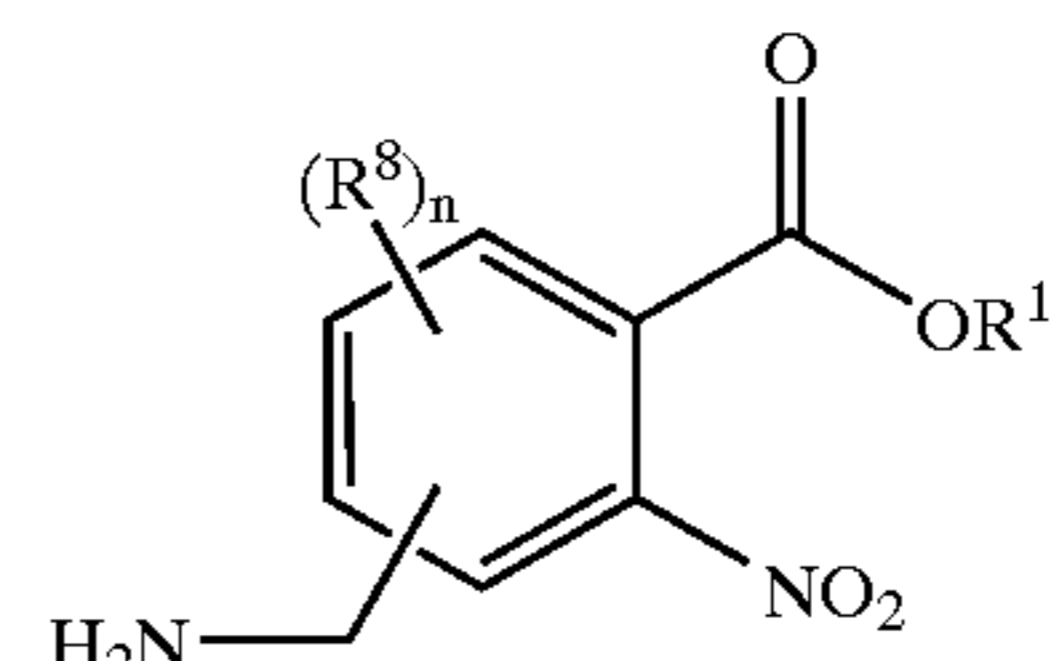
X is CH or N

and wherein, unless otherwise identified, the substituents are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₁-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C₁-C₈)

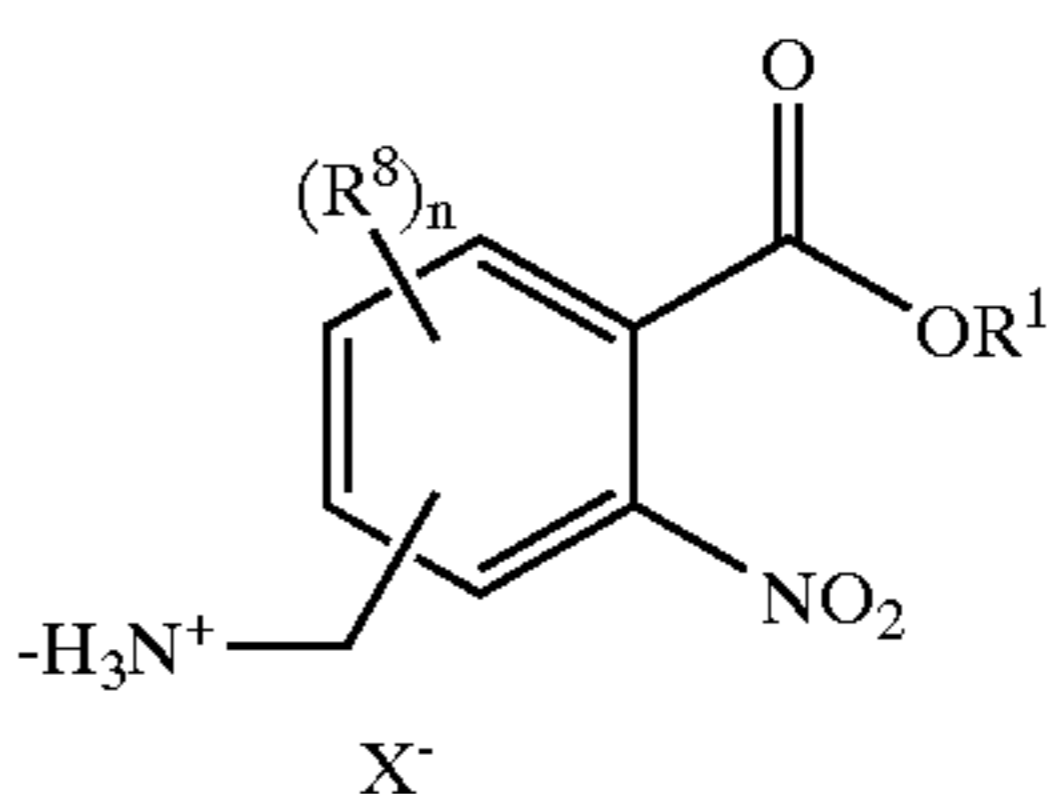
26

alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy; which comprises:

a) reacting a compound of the formulae

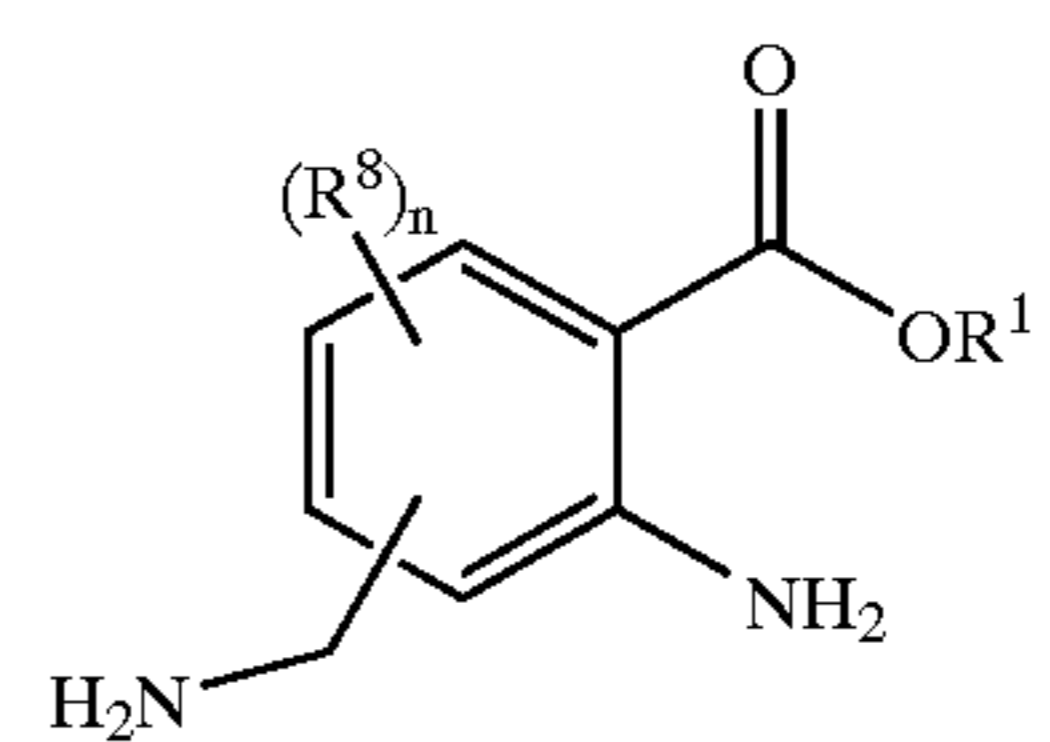


(V)

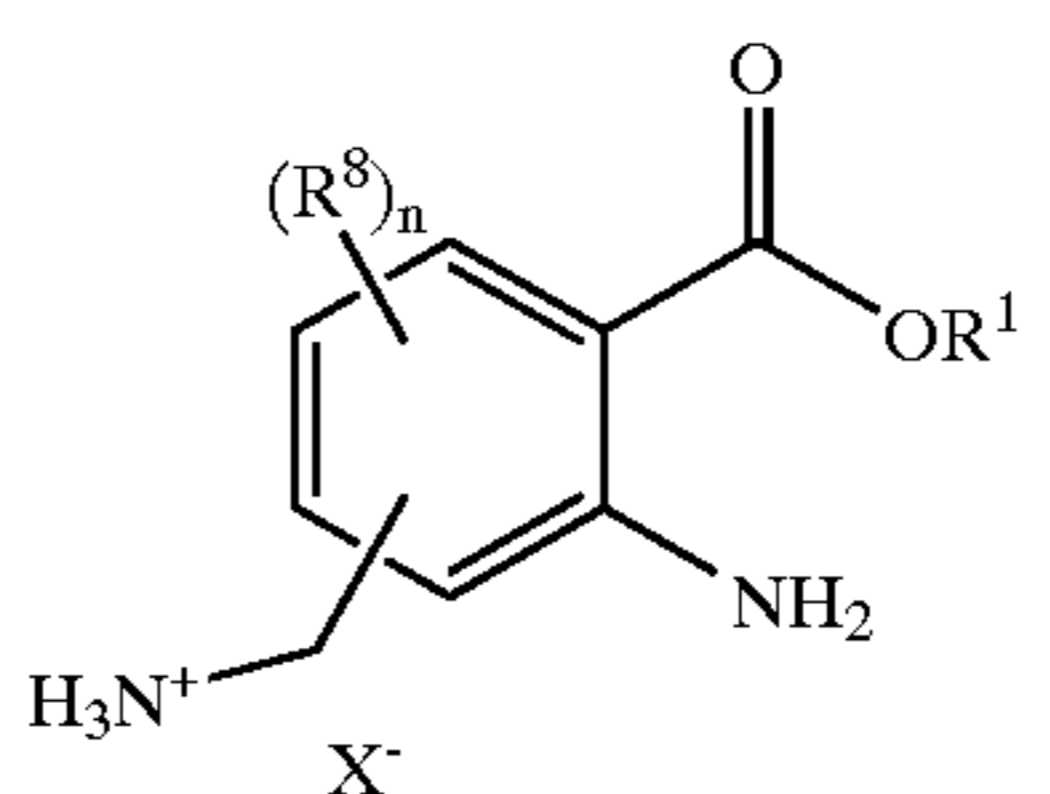


(Va)

with either a reagent that reduces a nitro group to an amine or by catalytic hydrogenation to give a compound of the formula

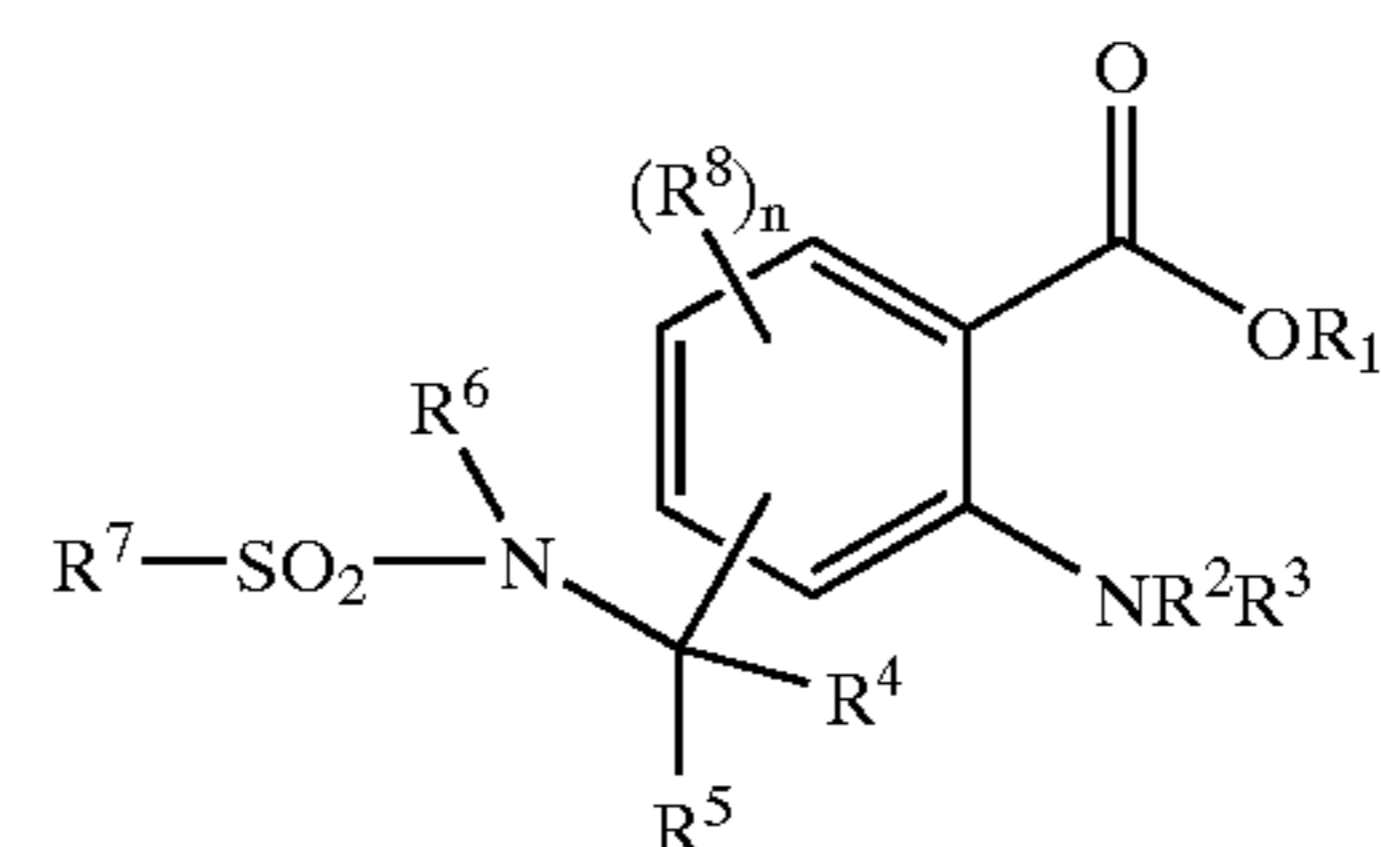


(III)



(IIIa)

b) reacting the compound of formula (III) or (IIIa) with a sulfonic acid derivative to give a compound of the formula



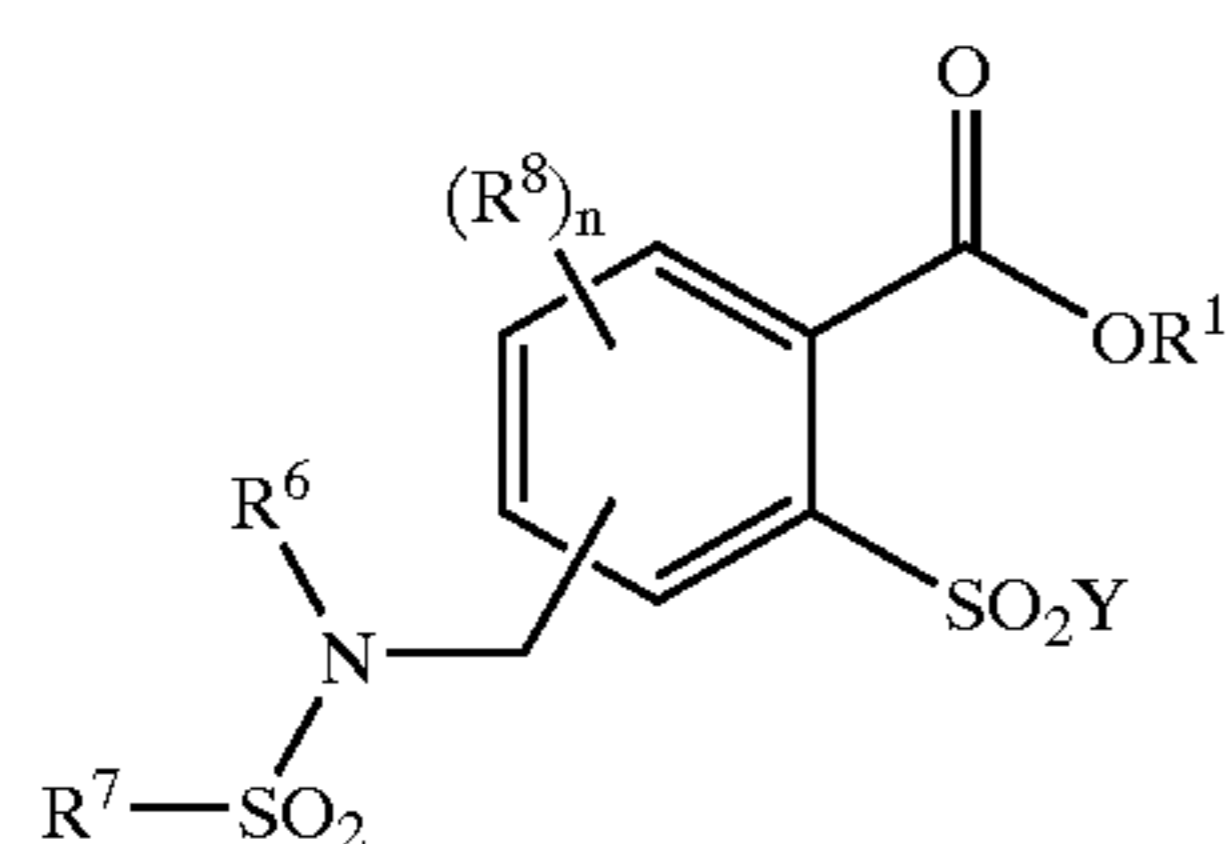
(I)

which R¹, R², and R⁶ are hydrogen

c) reacting the compound of formula (I) with an alkylating agent or by reductive animation to give a compound of the formula (I) where R⁶ is an unsubstituted or substituted C₁-C₈-alkyl group

d) first reacting the compound formed in step c) with a diazotizing agent in the presence of an acid followed by an SO₂ source in the presence of a copper catalyst and an acid to give a compound of the formula

27

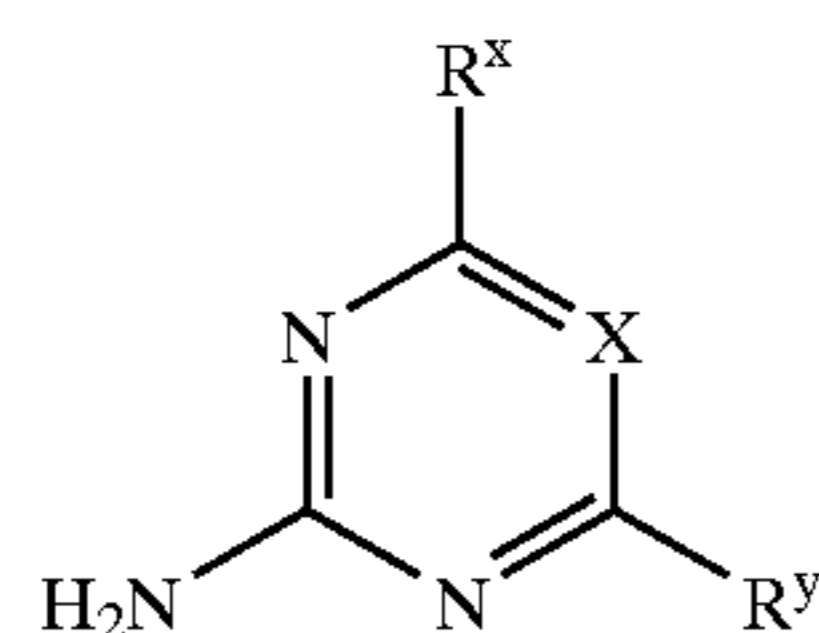


(VII)

where Y is halogen

e) reacting the compound of formula (VII) with either:

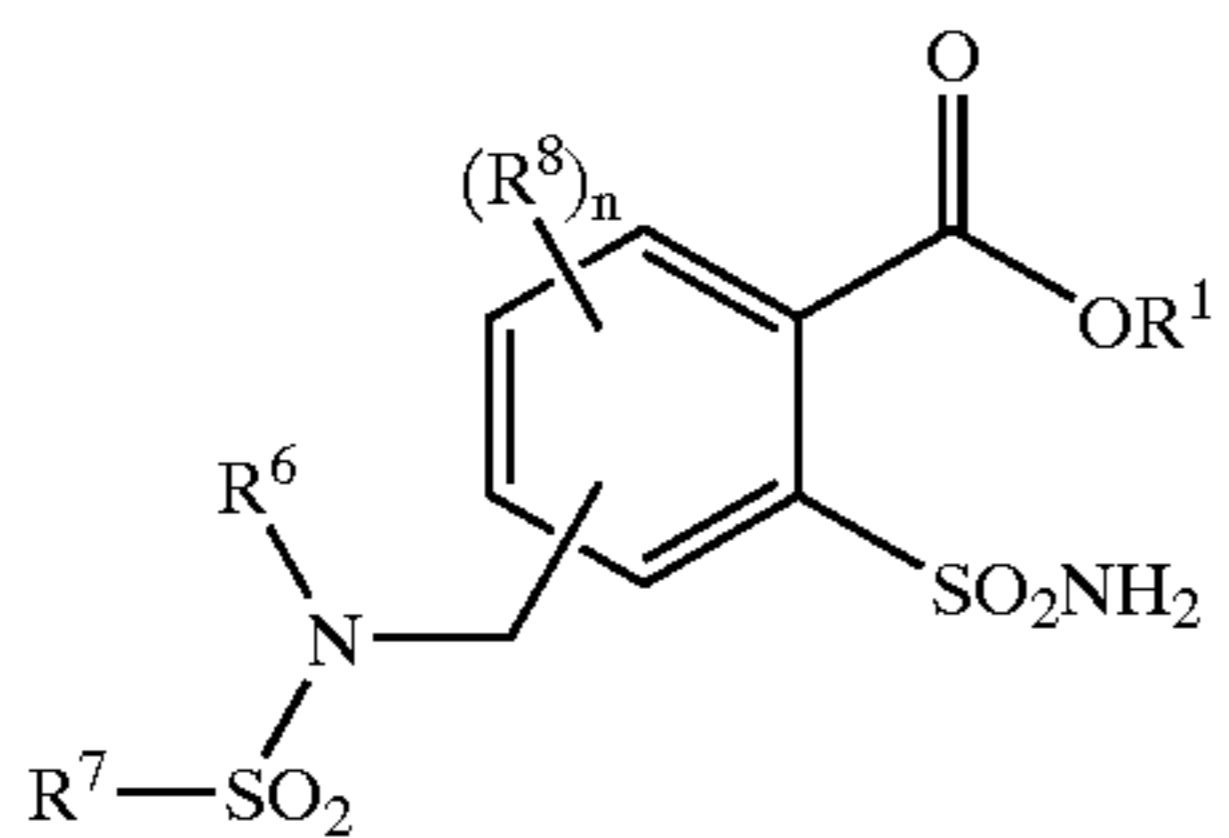
i) amine of the formula (XII) in the presence of MOCN, wherein M is an ammonium ion or an alkali metal ion, to give a compound of the formula (XIII)



(XII)

or

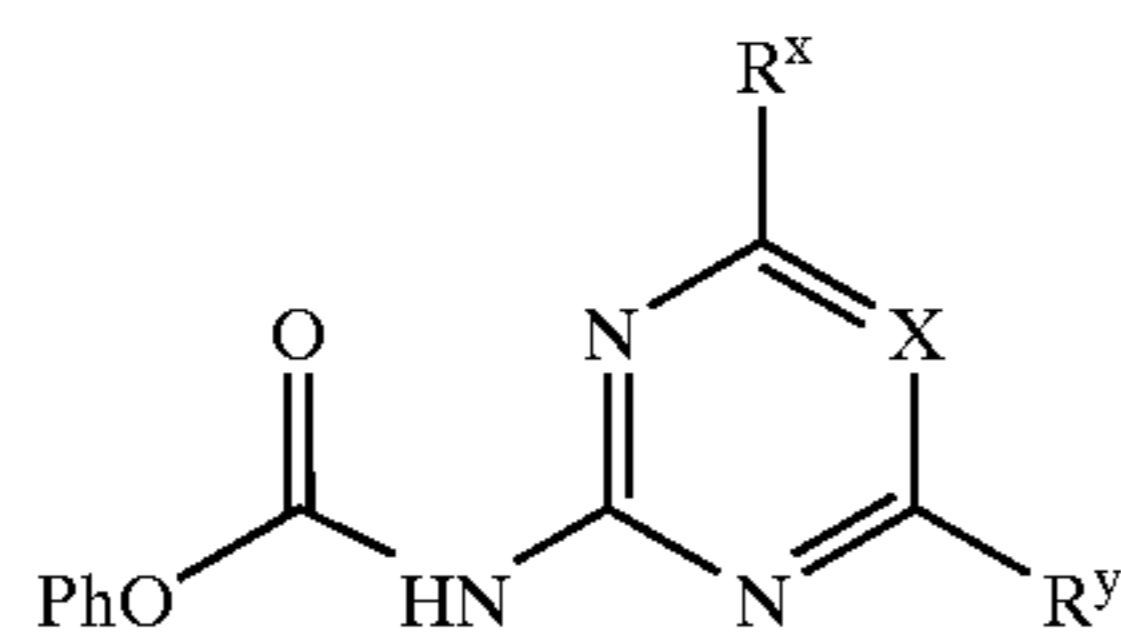
ii) ammonia in the presence of a suitable solvent to give a compound



(VIII)

and subsequently preparing the compound of the formula (XIII) by

a) reacting the compound of the formula (VIII) with a compound of the formula (IX) to give the compound of the formula (XIII)

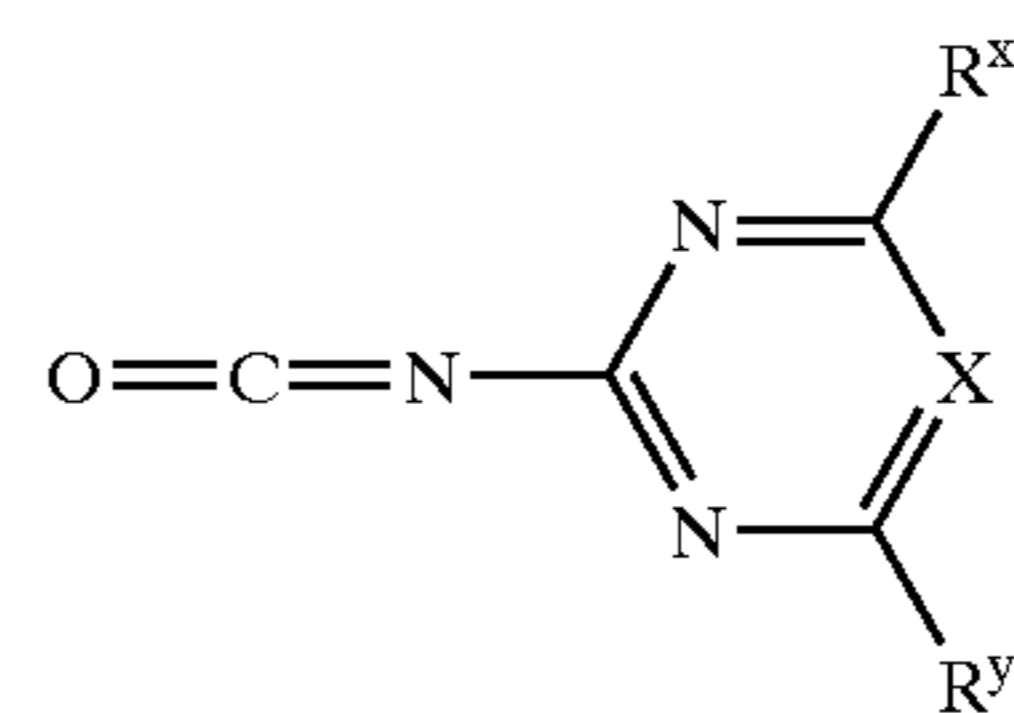


(IX)

or

b) reacting the compound of the formula (VIII) with an isocyanate of the formula (X) to give the compound of the formula (XIII)

28



(X)

5

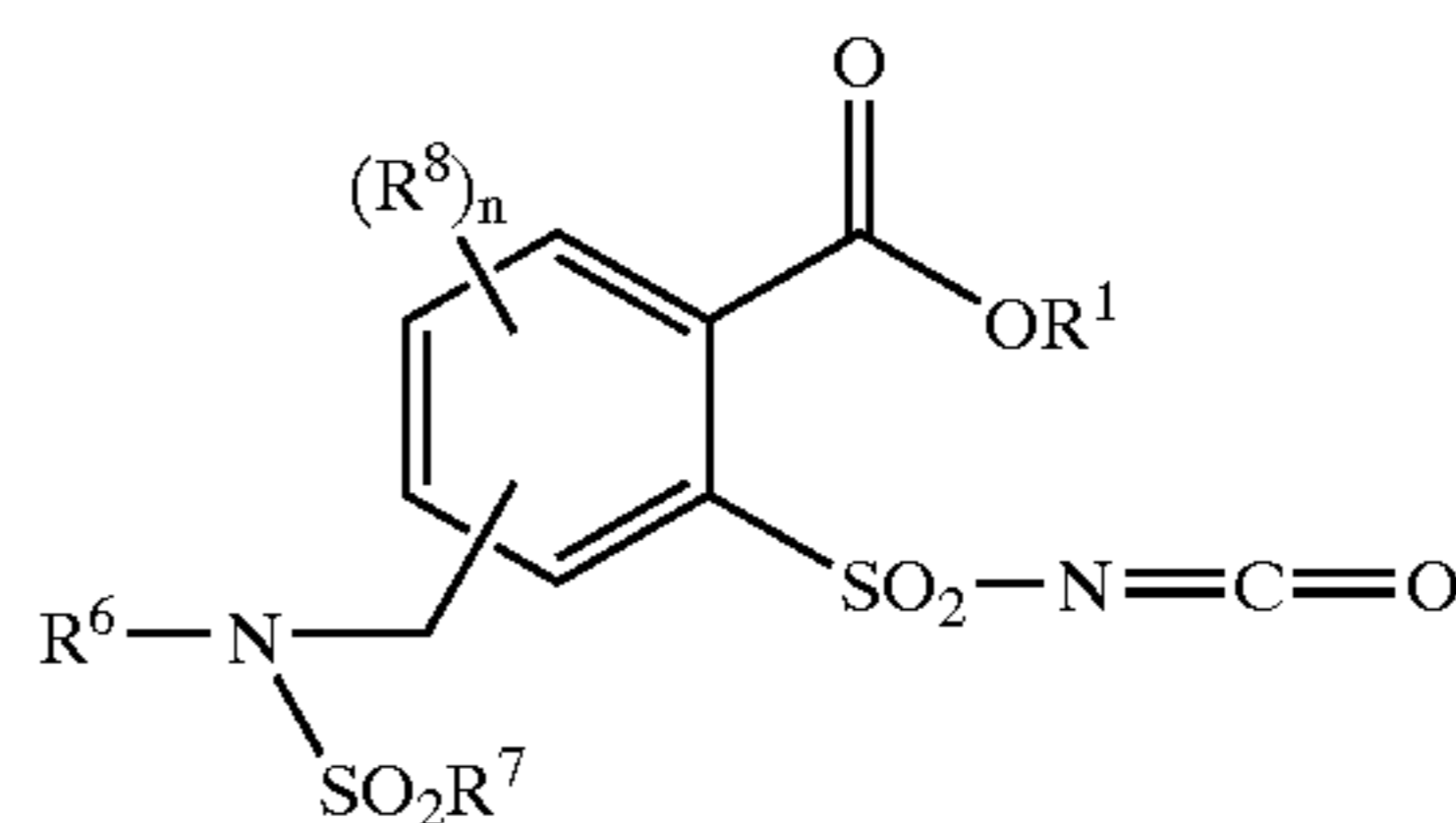
10

or

c) reacting the compound of the formula (VIII) with an alkyl isocyanate and phosgene to give a compound of the formula (XI)

15

(XI)



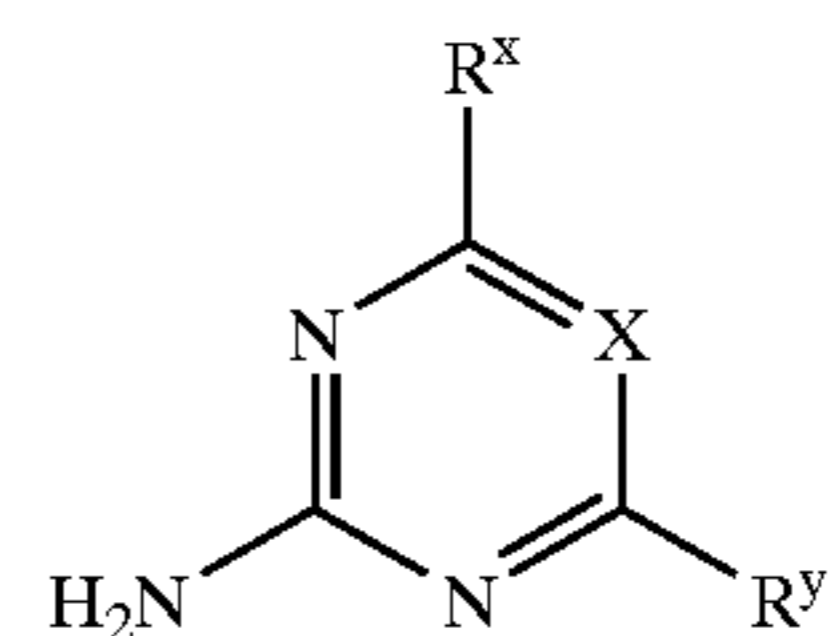
20

25

which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

(XII)

30



35

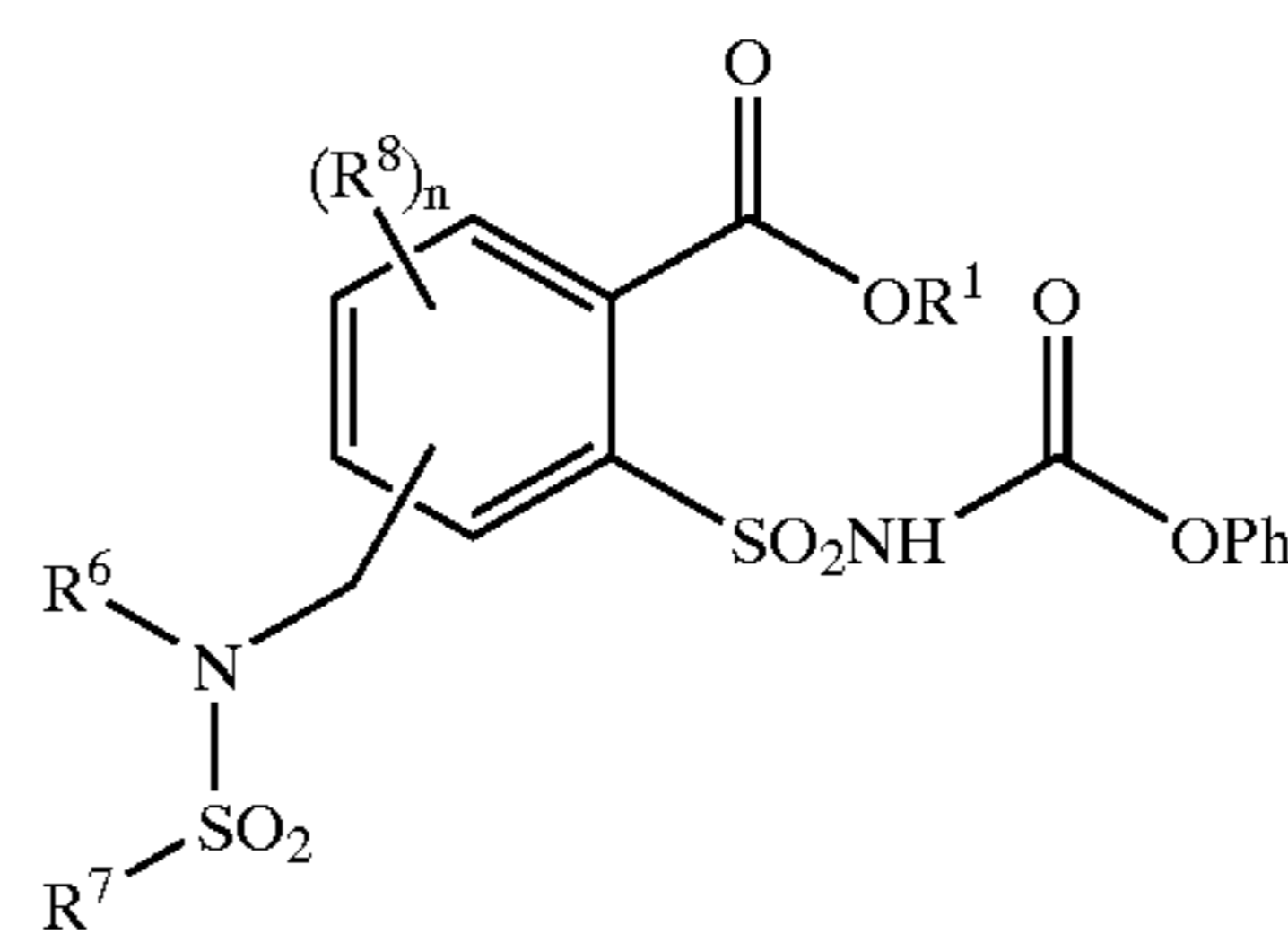
or

d) reacting the compound of the formula (VIII) with a carbonic acid derivative R-CO-OPh, in which Ph=unsubstituted or substituted phenyl and R=halogen or unsubstituted or substituted phenoxy to give a compound of the formula (XIV)

40

(XIV)

45

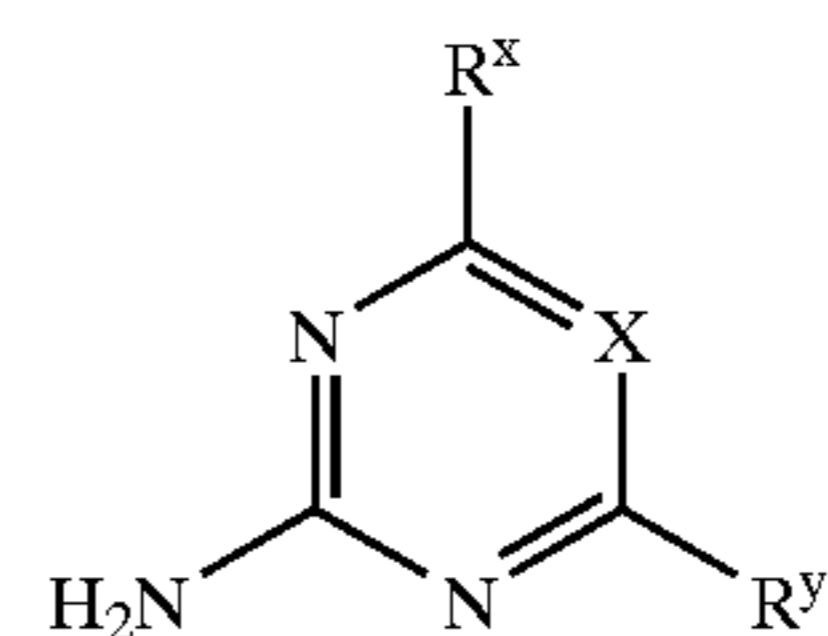


50

(IX) which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

(XII)

60



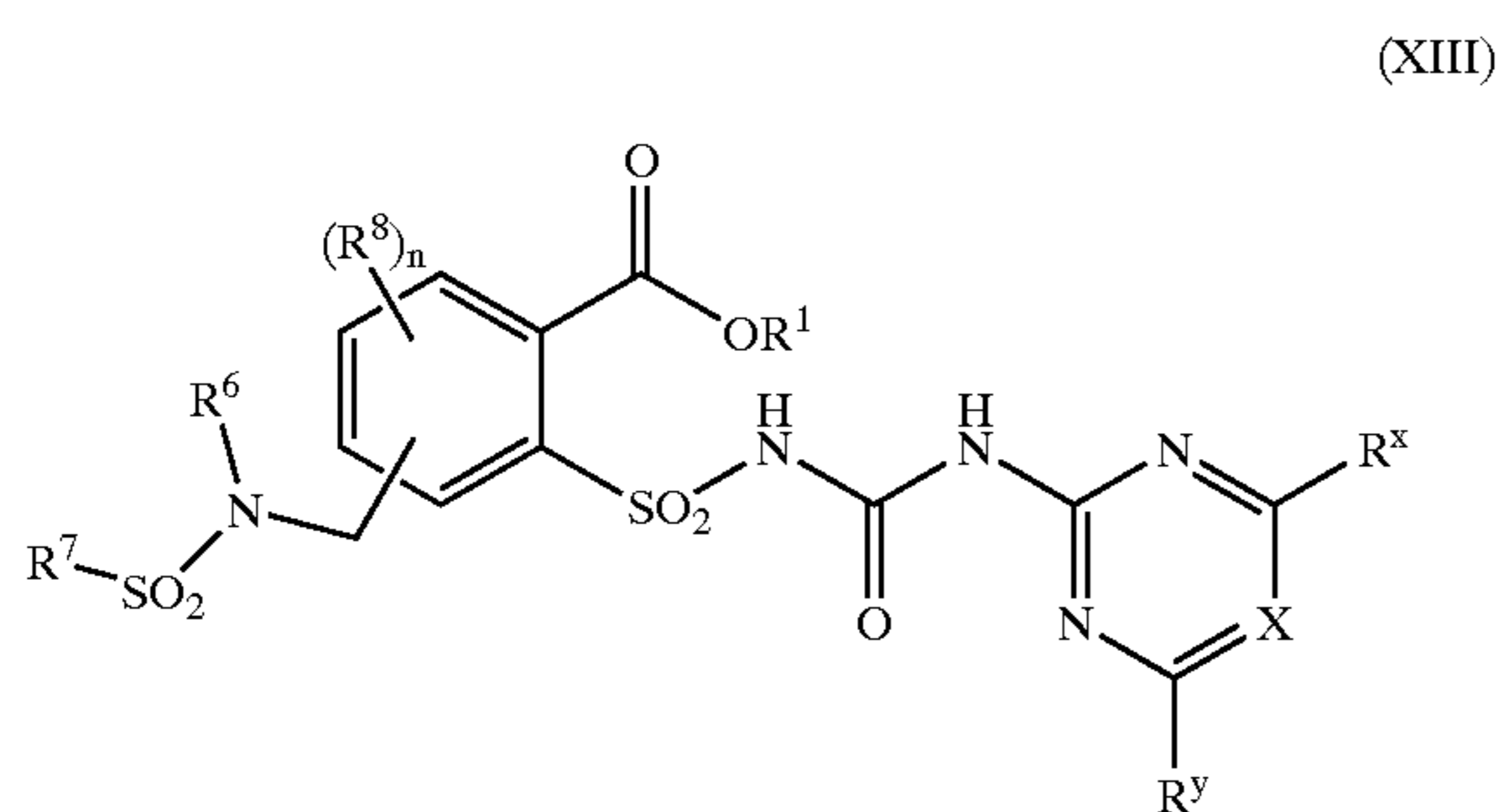
where R¹, R⁷, R⁸, and n in formulae (XI) and (XIV) are as defined in formula (1) R^x, R^y and X in formulae (IX), (X)

65

29

and (XII) are as defined in formula (XIII) of the present claim and Ph in formulae (IX) and (XIV) is unsubstituted or substituted phenyl, wherein the substituents on the phenyl ring are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₃-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C₁-C₈)alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy.

4. A process for preparing a phenylsulfonylurea of the formula



in which

R¹ is H, (C₁-C₈)alkyl, (C₃-C₈)alkenyl or (C₃-C₈)alkynyl, where the last 3 radicals are unsubstituted or substituted,

R⁶ is H or (C₁-C₈)alkyl which is unsubstituted or substituted,

R⁷ is (C₁-C₈)alkyl, (C₃-C₈)alkenyl, (C₃-C₈)alkynyl, (C₆-C₁₄)aryl or mono- or di(C₁-C₈)alkylamino which are unsubstituted or substituted, or

R⁶ and R⁷ together form a chain of the formula —(CH₂)_m E_m¹— which is unsubstituted or substituted, and where m=2, 3 or 4, m¹=0 or 1 and E=CO or SO₂

R⁸ radicals are identical or different and are (C₁-C₄)alkyl, (C₁-C₄)alkoxy, [(C₁-C₄)alkyl]carbonyl or [(C₁-C₄)alkoxy]carbonyl which are unsubstituted or substituted, or R⁸ is halogen, NO₂ or CN, and

n is 0, 1, 2 or 3,

R^x, R^y independently of one another are a hydrogen atom, halogen, (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)alkylthio, where each of the last-mentioned 3 radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, (C₁-C₄)alkoxy and (C₁-C₄)alkylthio, or are mono- or di[(C₁-C₄)alkyl]amino, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, (C₃-C₆)alkenyloxy or (C₃-C₆)alkynyloxy, and

X is CH or N

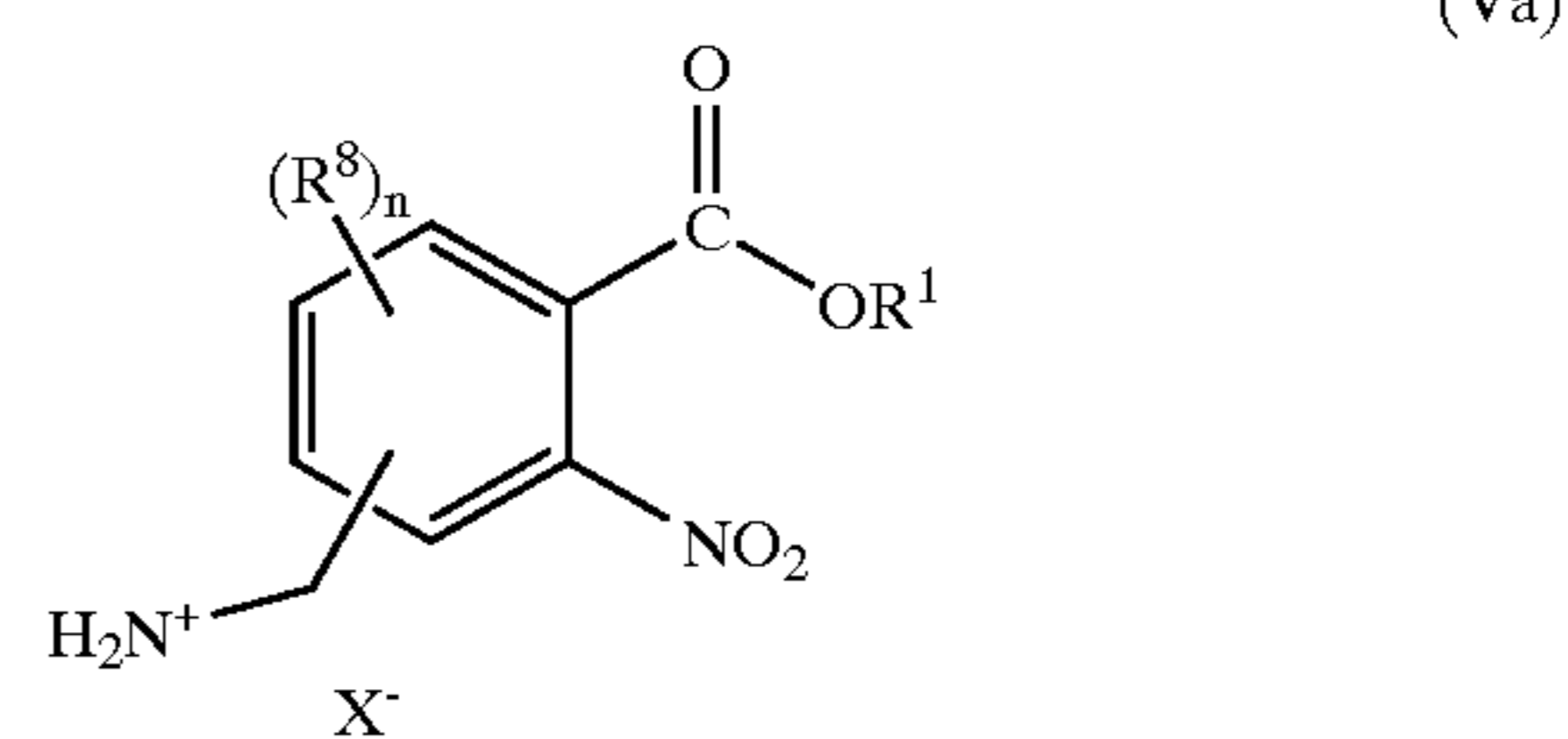
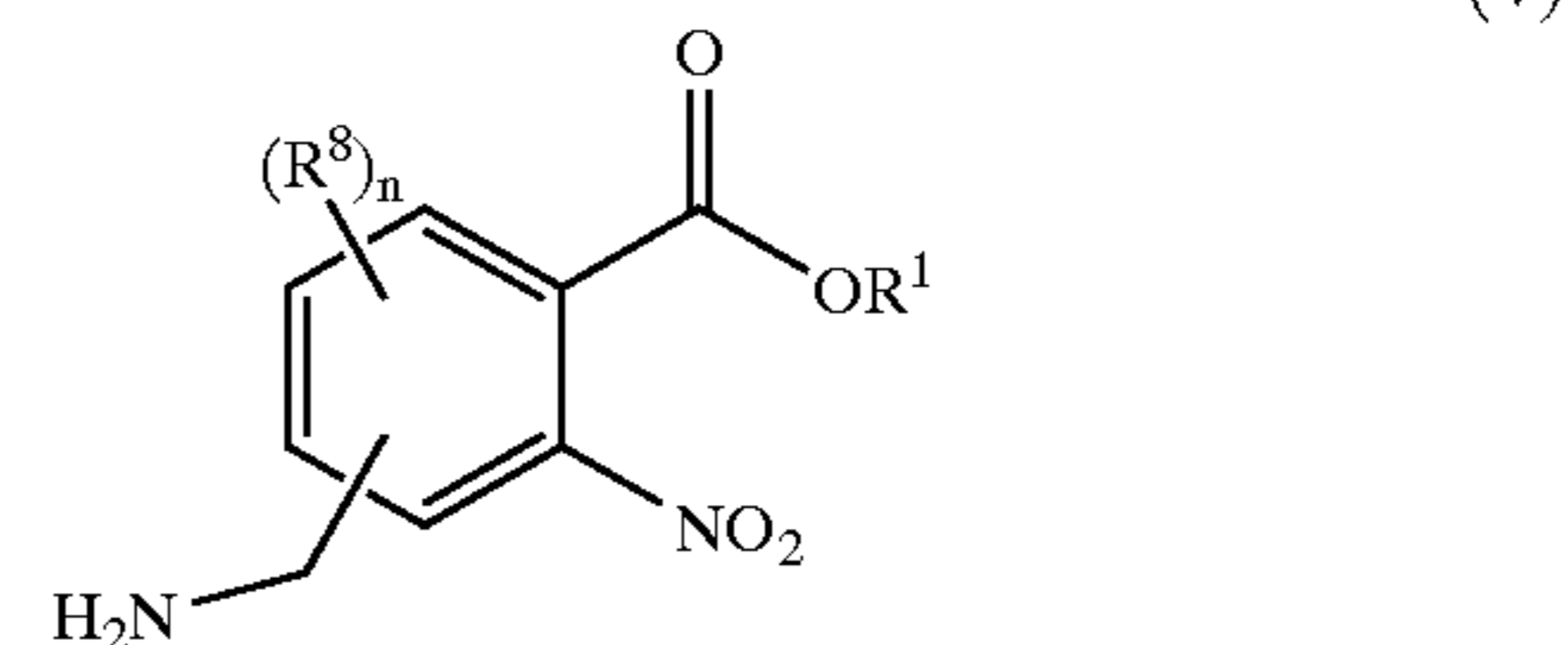
and wherein, unless otherwise identified, the substituents are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₁-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl

30

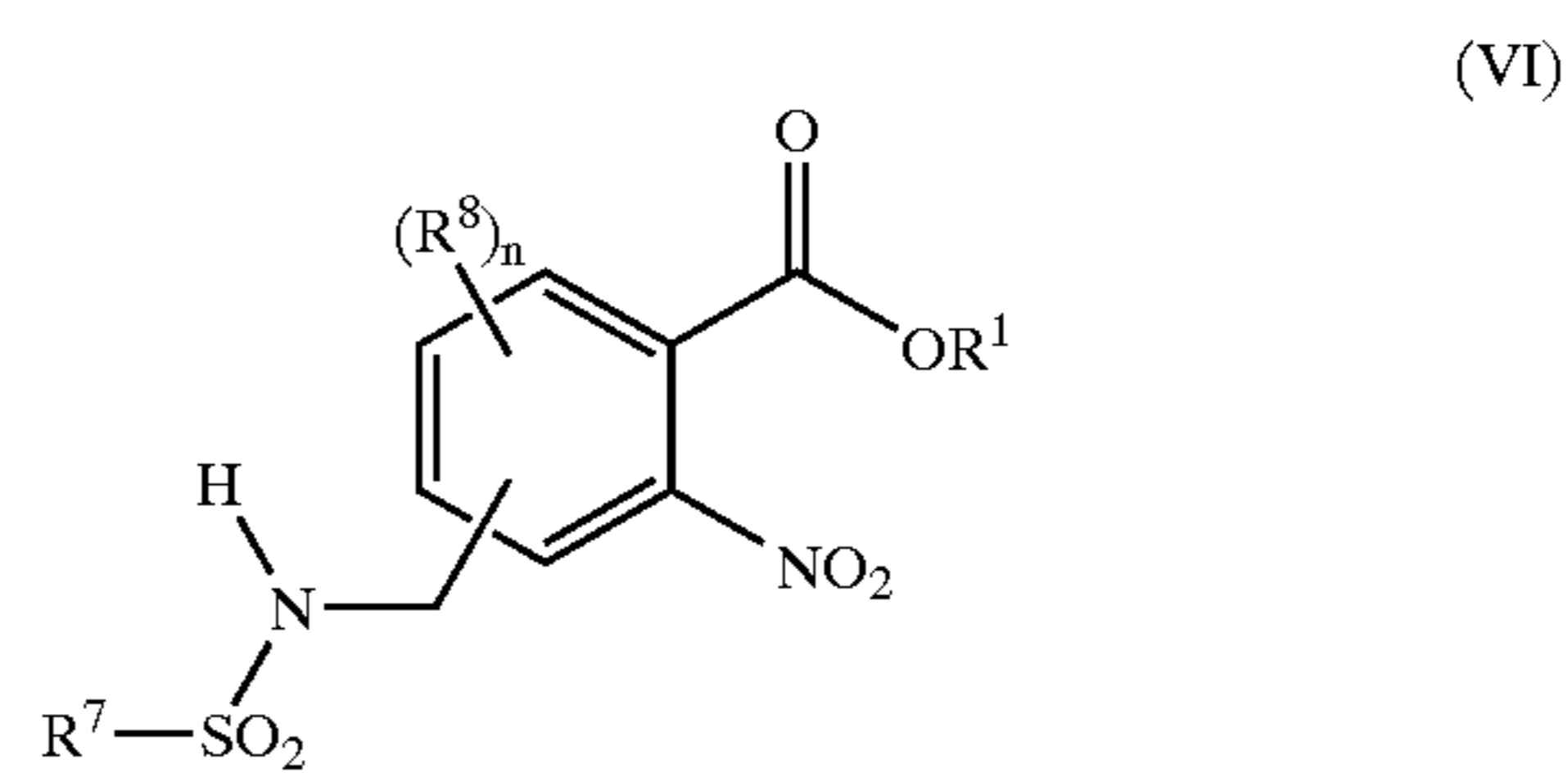
and cycloalkyl radicals are optionally substituted by (C₁-C₈)alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy,

which comprises:

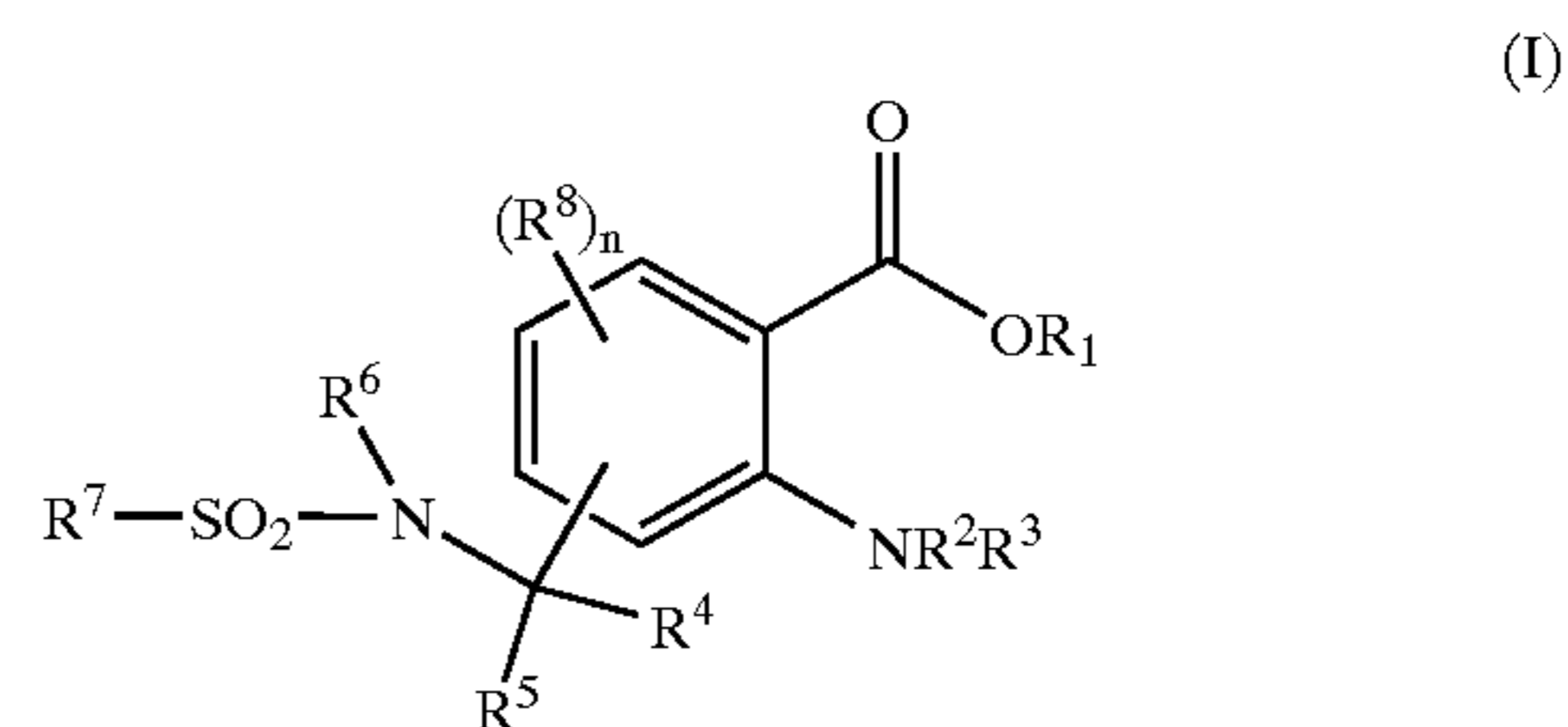
a) reacting a compound of formula



with a sulfonic acid derivative to give a compound of the formula



and reacting the compound of the formula (VI) with a reagent that reduces a nitro group to an amine or by catalytic hydrogenation to give a compound of the formula

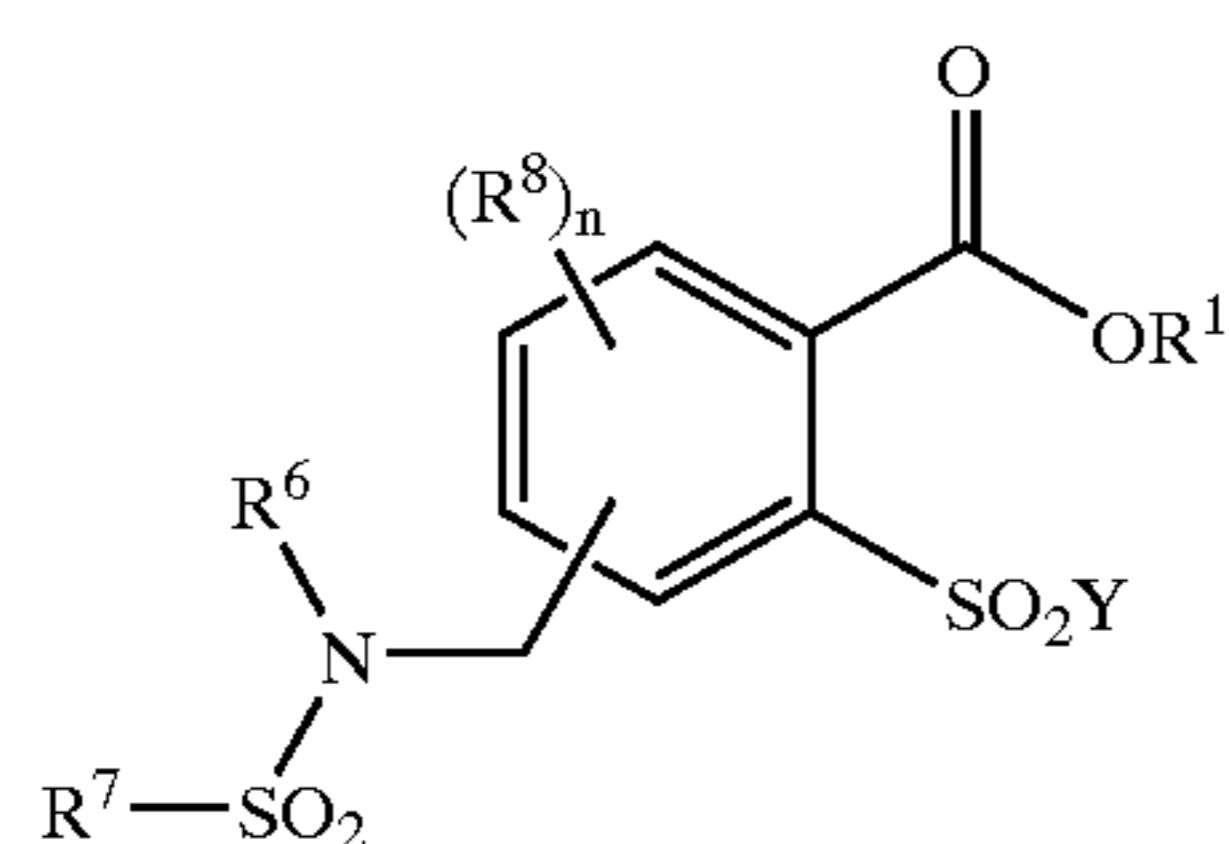


which R², R³, and R⁶ are hydrogen

b) reacting the compound of formula (I) with an alkylating agent or by reductive amination to give a compound of the formula (I) where R⁶ is an unsubstituted or substituted C₁-C₈-alkyl group

c) first reacting the compound formed in step b) with a diazotizing agent in the presence of an acid followed by an SO₂ source in the presence of a copper catalyst and an acid to give a compound of the formula

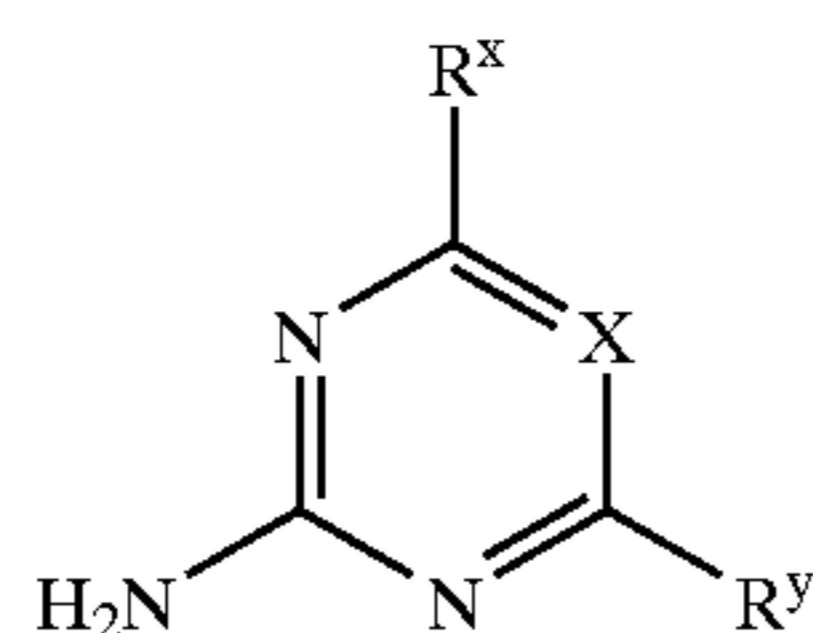
31



where Y is halogen

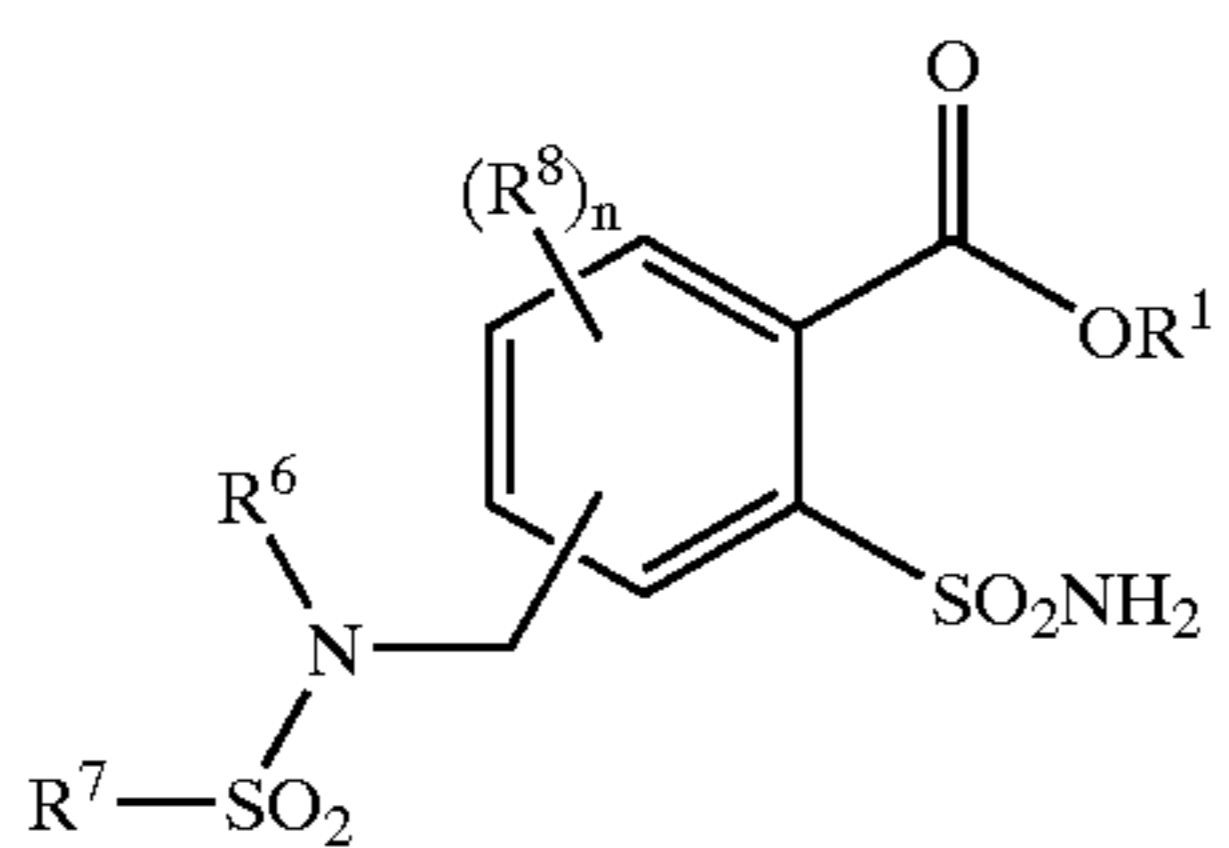
d) reacting the compound of formula (VII) with either:

i) amine of the formula (XII) in the presence of MOCN, wherein M is an ammonium ion or an alkali metal ion, to give a compound of the formula (XIII)



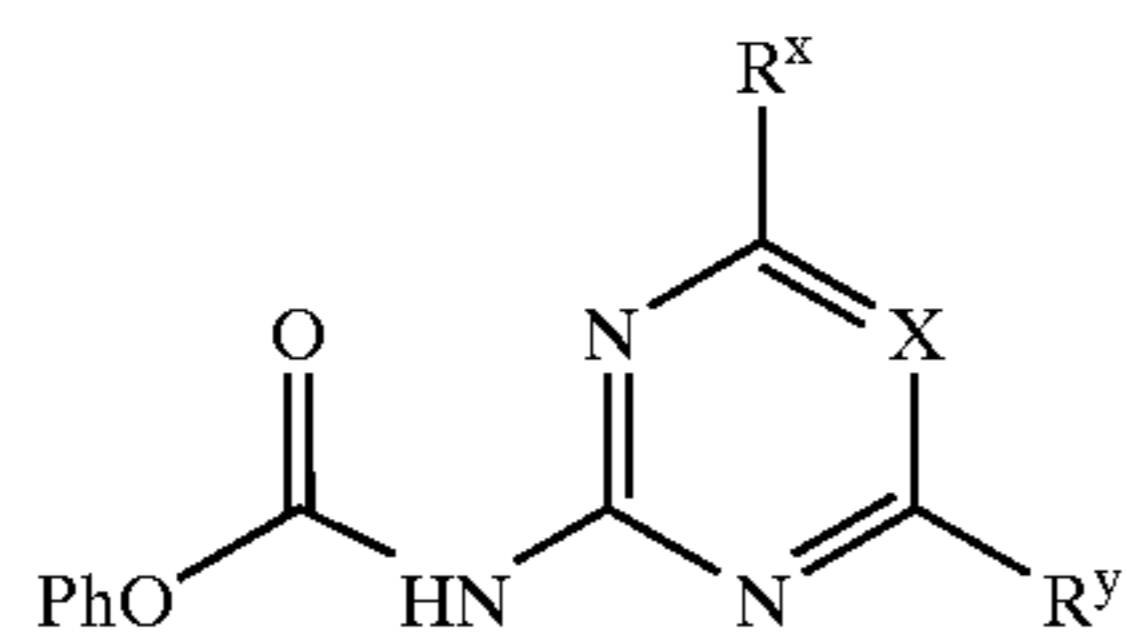
or

ii) ammonia in the presence of a suitable solvent to give a compound



and subsequently preparing the compound of the formula (XIII) by:

a) reacting the compound of the formula (VIII) with a compound of the formula (IX) to give the compound of the formula (XIII)



or

b) reacting the compound of the formula (VIII) with an isocyanate of the formula (X) to give the compound of the formula (XIII)

(VII)

5

10

15

20

(XIII)

25

30

(VIII)

40

45

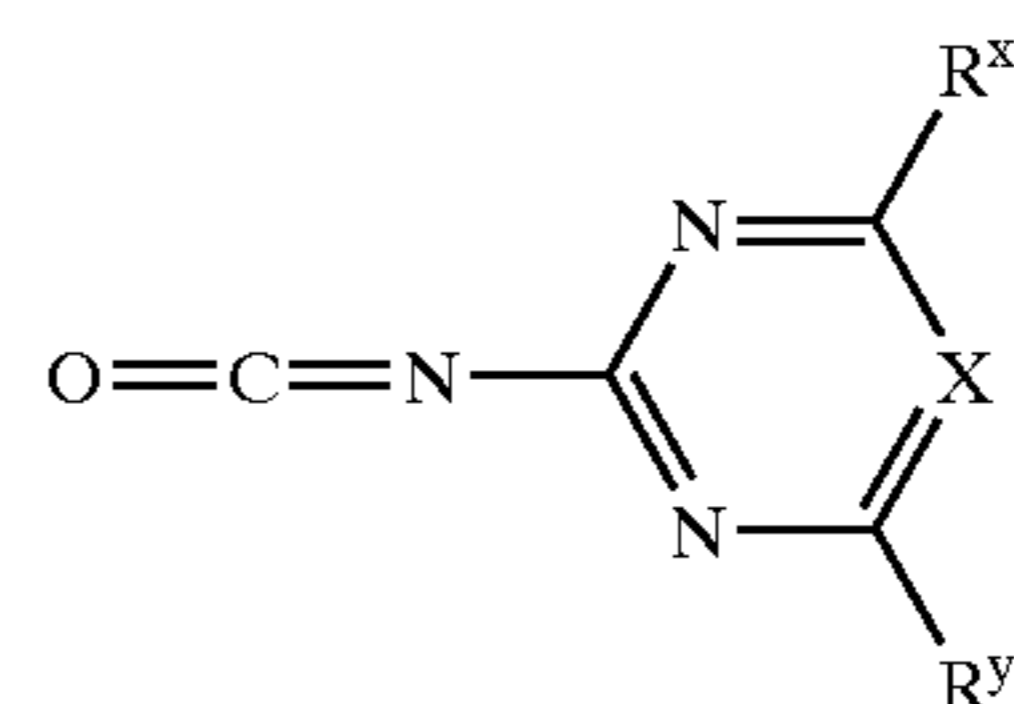
50

(IX)

60

65

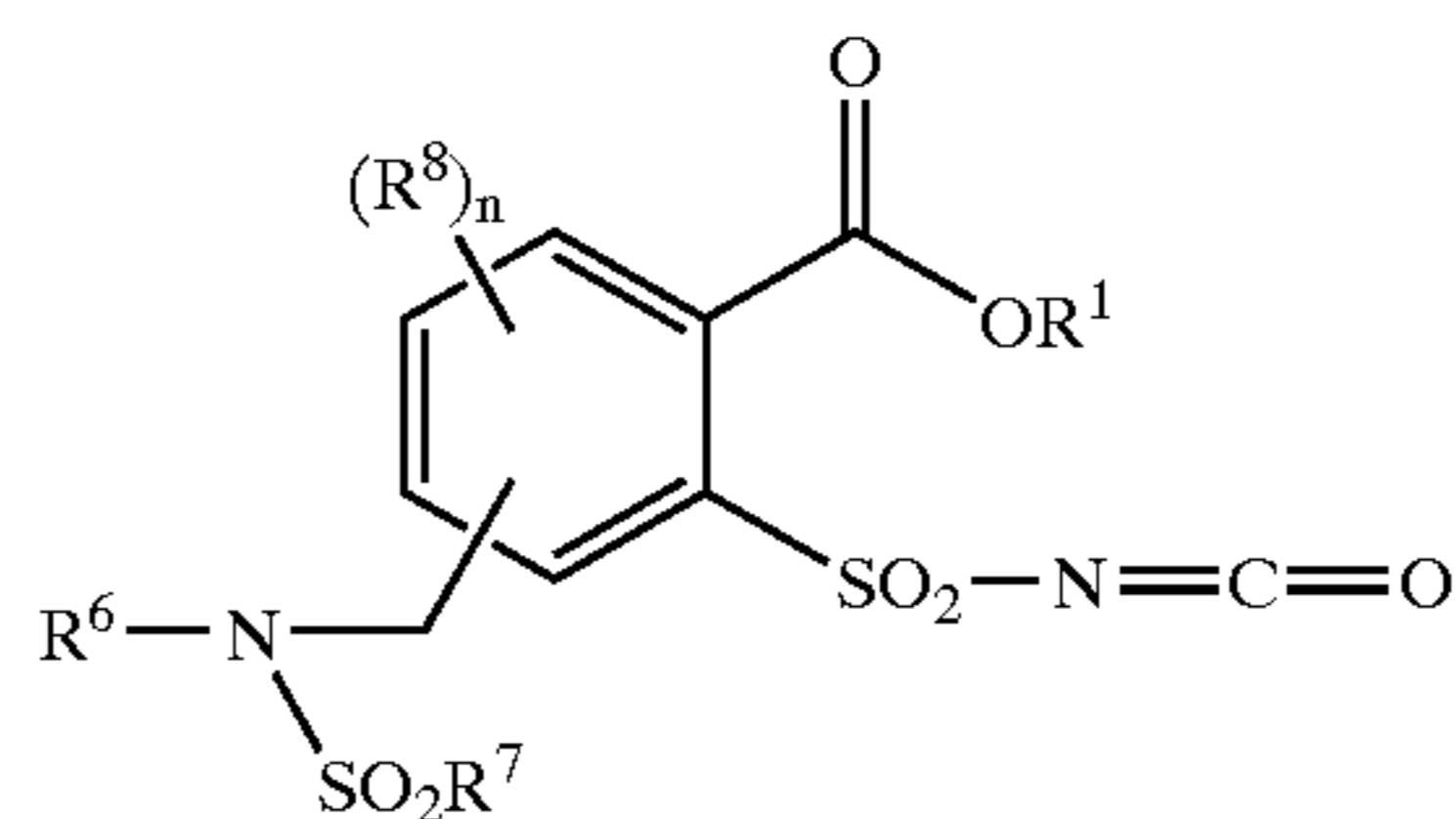
32



(X)

or

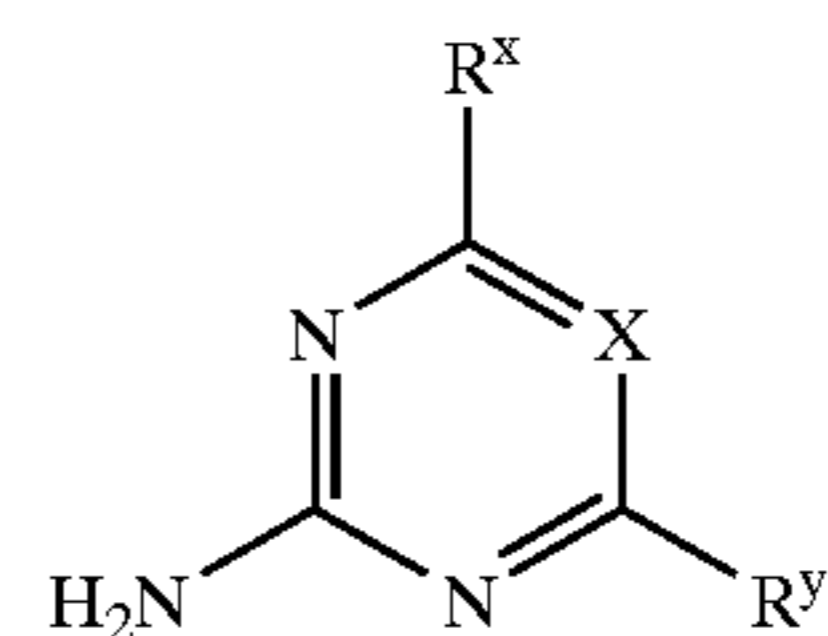
c) reacting the compound of the formula (VIII) with an alkyl isocyanate and phosgene to give a compound of the formula (XI)



(XI)

which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

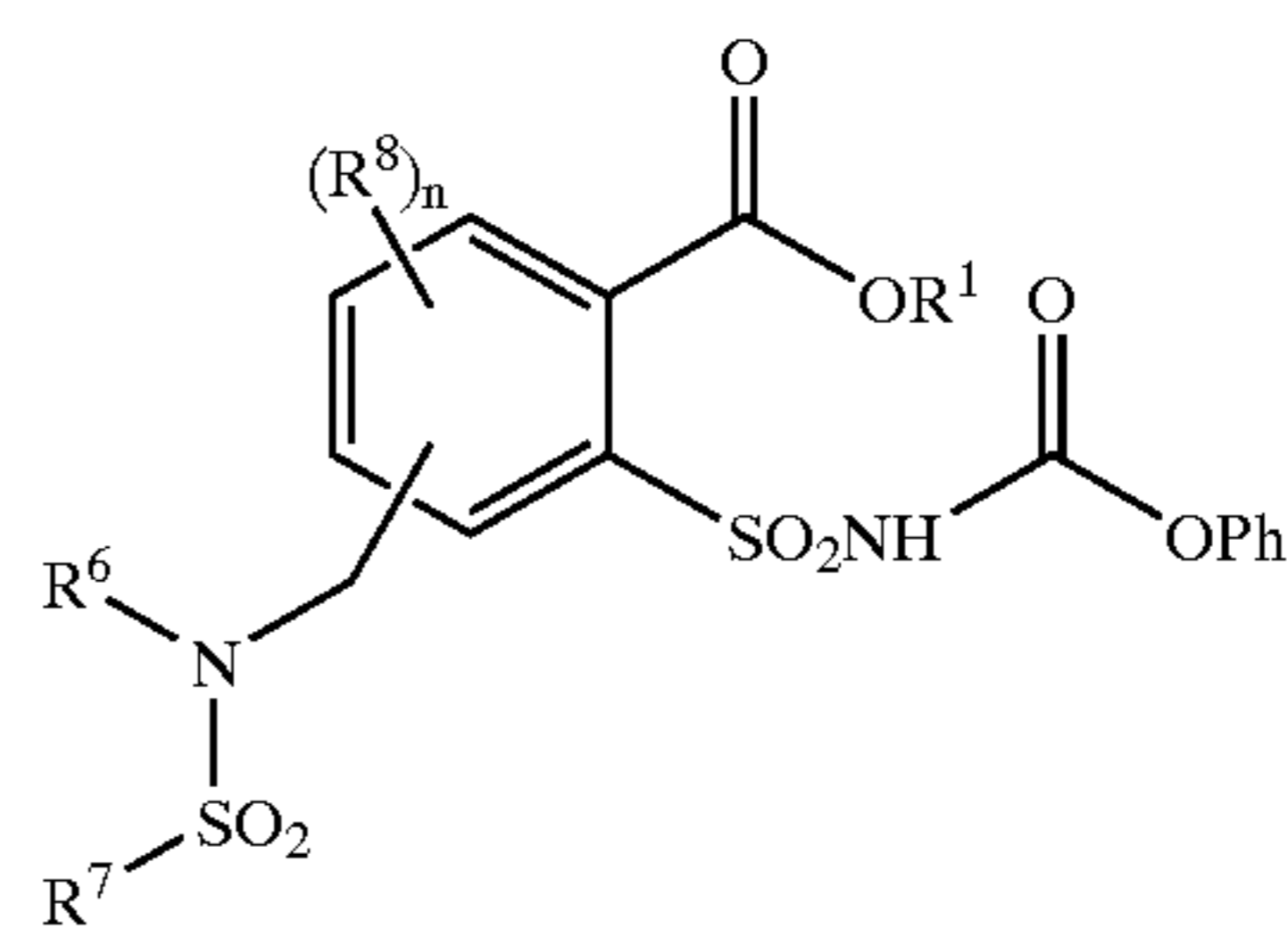
(XII)



or

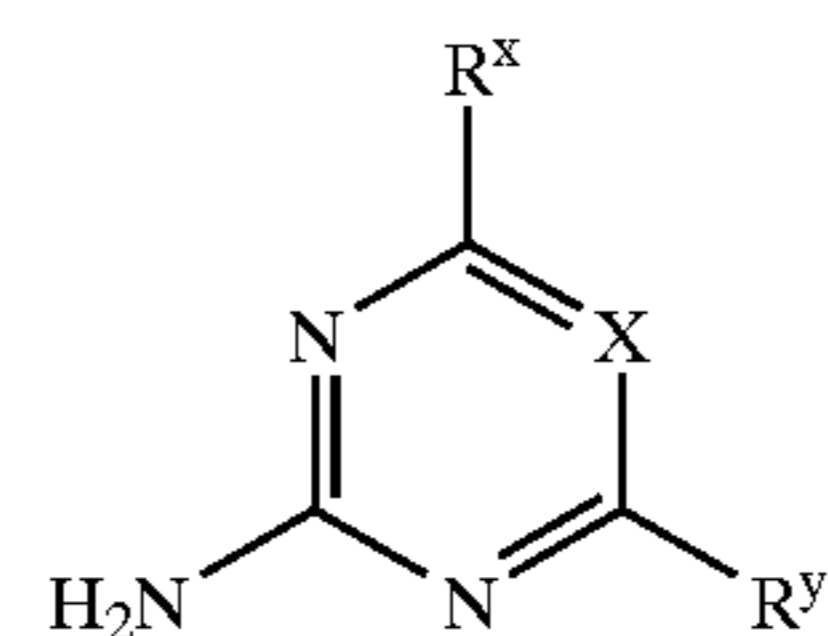
d) reacting the compound of the formula (VIII) with a carbonic acid derivative R-CO-OPh, in which Ph=unsubstituted or substituted phenyl and R=halogen or unsubstituted or substituted phenoxy to give a compound of the formula (XIV)

(XIV)



which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

(XII)

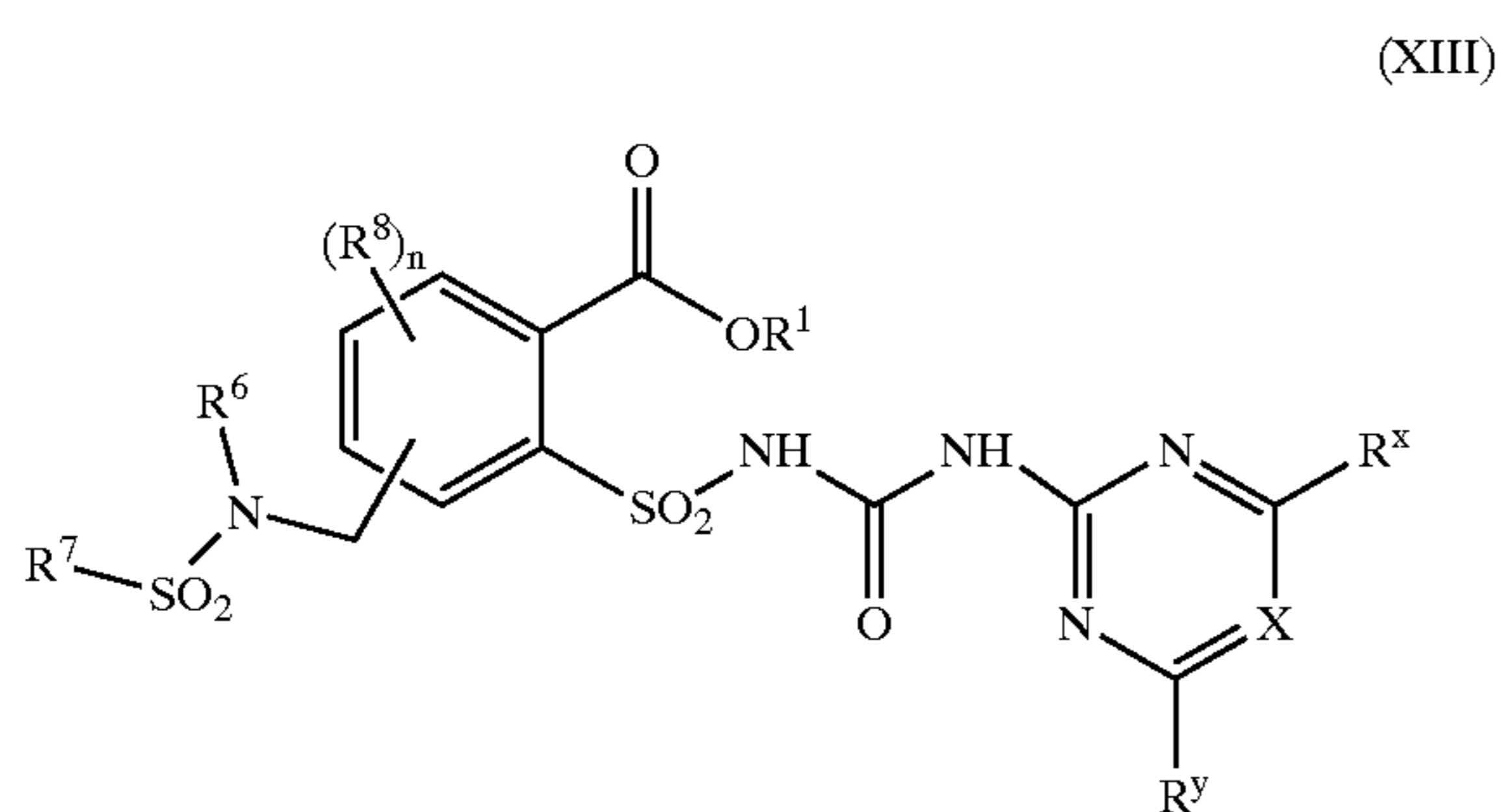


where R¹, R⁷, R⁸, and n in formulae (XI) and (XIV) are as defined in formula (I), R^x, R^y and X in formulae (IX), (X)

33

and (XII) are as defined in formula (XIII) of the present claim and Ph in formulae (IX) and (XIV) is unsubstituted or substituted phenyl, wherein the substituents on the phenyl ring are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₁-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C₁-C₈)alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy.

5. A process for preparing a phenylsulfonylurea of the formula



in which

R¹ is H, (C₁-C₈)alkyl, (C₃-C₈)alkenyl or (C₃-C₈)alkynyl, where the last 3 radicals are unsubstituted or substituted,

R⁶ is H or (C₁-C₈)alkyl which is unsubstituted or substituted,

R⁷ is (C₁-C₈)alkyl, (C₃-C₈)alkenyl, (C₃-C₈)alkynyl, (C₆-C₁₄)aryl or mono- or di(C₁-C₈)alkylamino which are unsubstituted or substituted, or

R⁶ and R⁷ form chain of the formula $-(CH_2)_m E m^1-$ which is unsubstituted or substituted, and where m=2, 3 or 4, m¹=0 or 1 and E=CO or SO₂,

R⁸ radicals are identical or different and are (C₁-C₄)alkyl, (C₁-C₄)alkoxy, [(C₁-C₄)alkyl]carbonyl or [(C₁-C₄)alkoxy]carbonyl which are unsubstituted or substituted, or R⁸ is halogen, NO₂ or CN, and

n is 0, 1, 2 or 3,

R^x, R^y independently of one another are a hydrogen atom, halogen, (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)alkylthio, where each of the last-mentioned 3 radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, (C₁-C₄)alkoxy and (C₁-C₄)alkylthio, or are mono- or di[(C₁-C₄)alkyl]amino, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, (C₃-C₆)alkenyloxy or (C₃-C₆)alkynyloxy, and

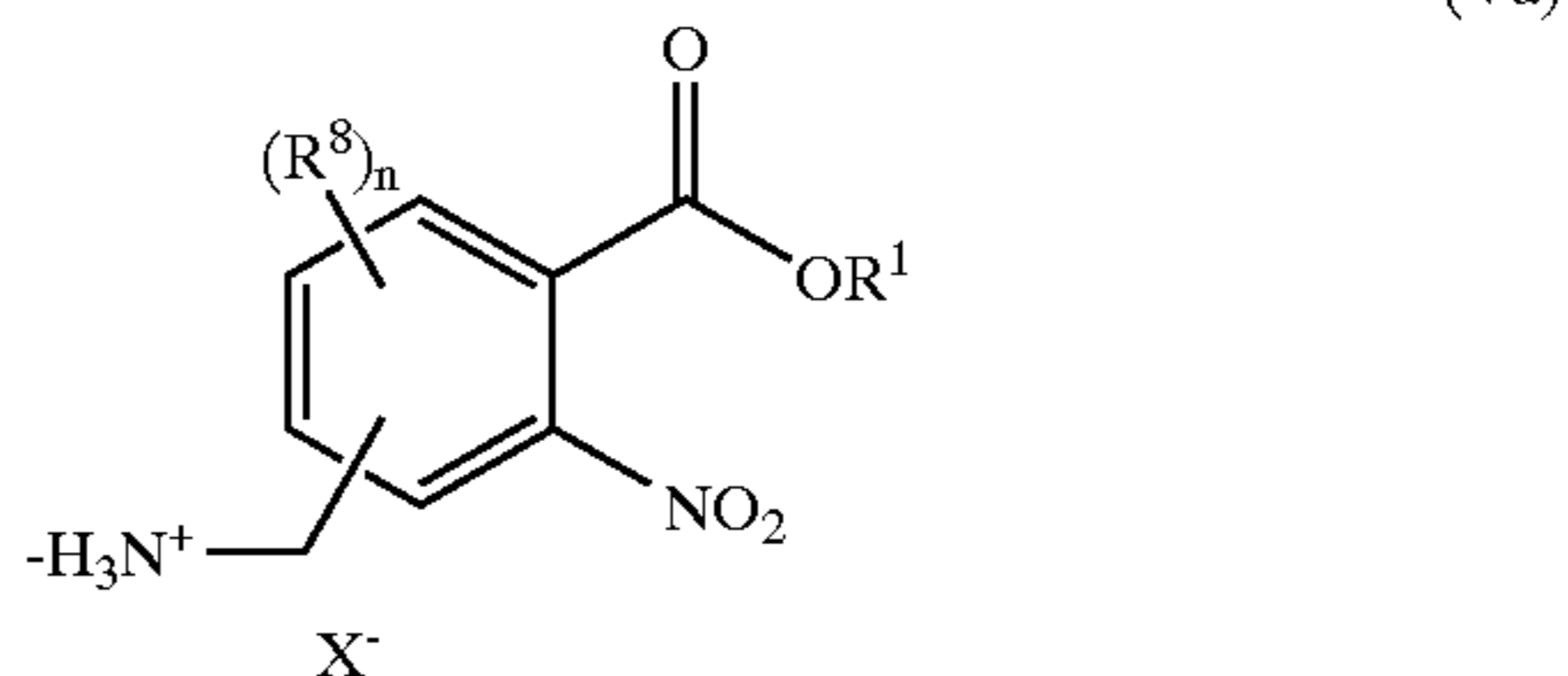
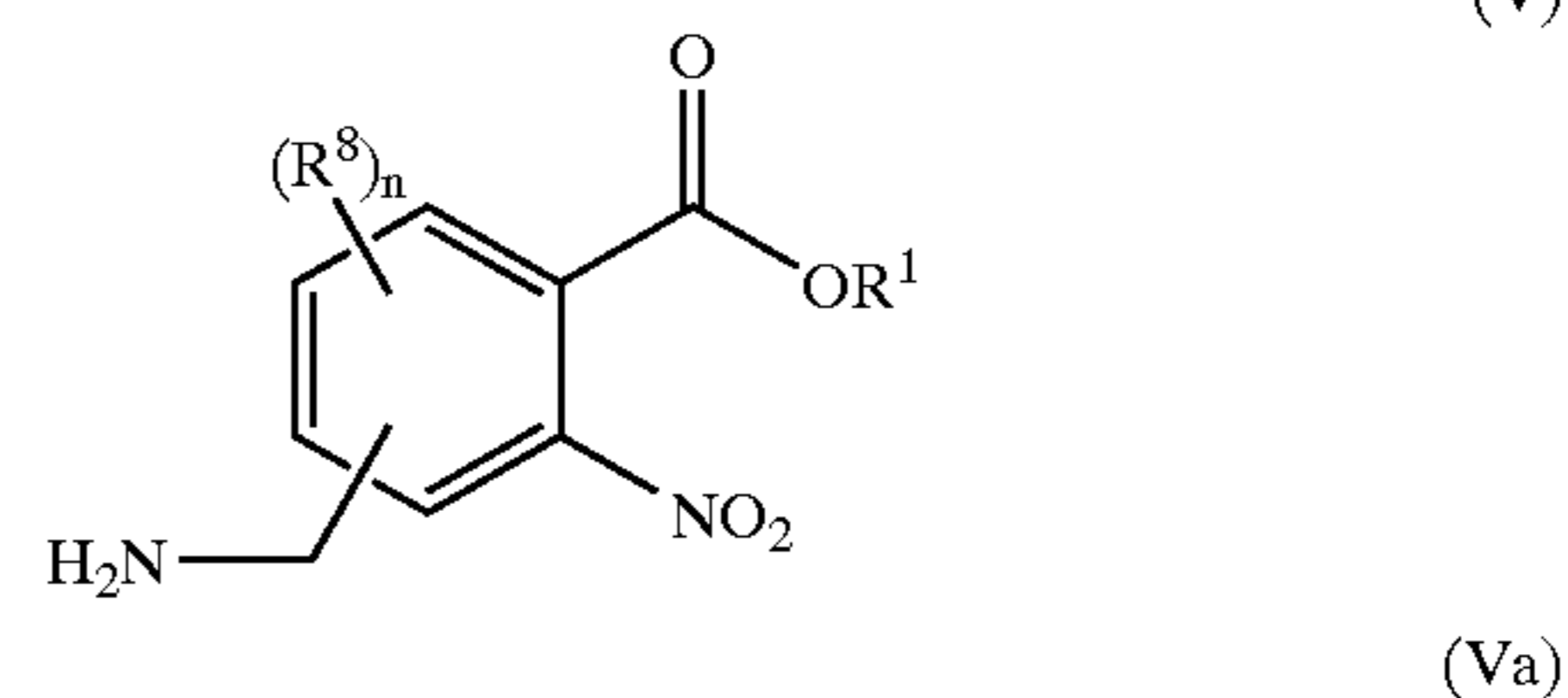
X is CH or N

and wherein, unless otherwise identified, the substituents are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₁-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl

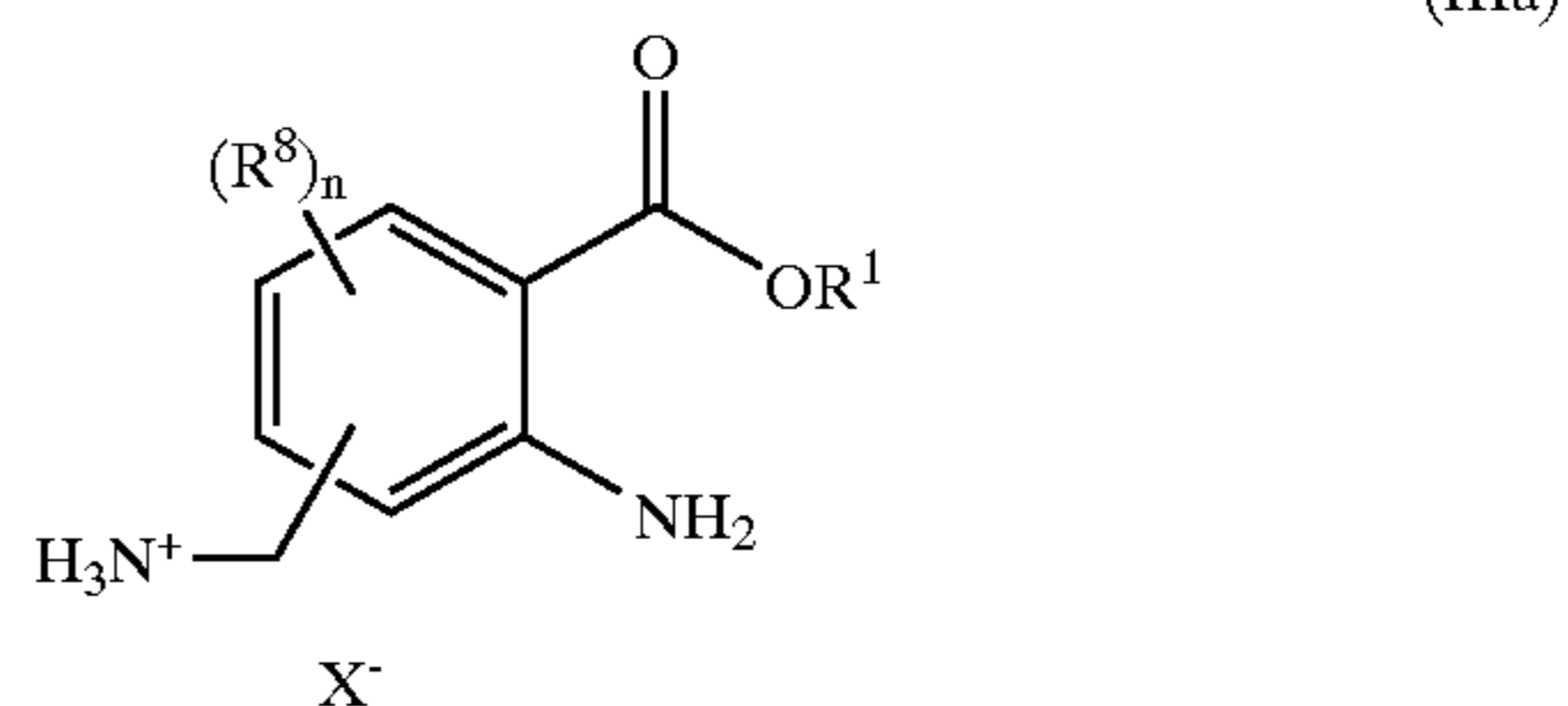
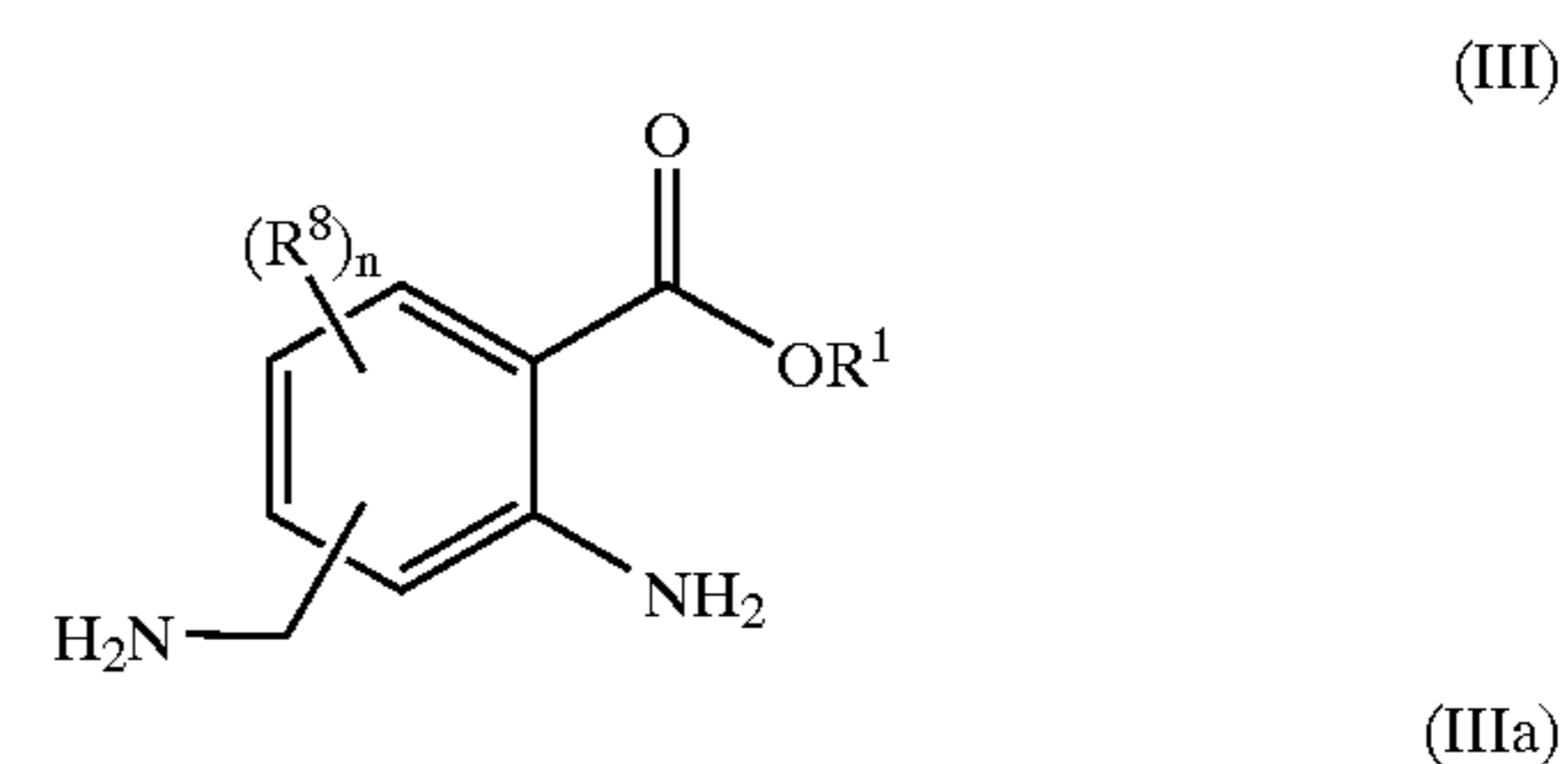
34

and cycloalkyl radicals are optionally substituted by (C₁-C₈)alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy, which comprises:

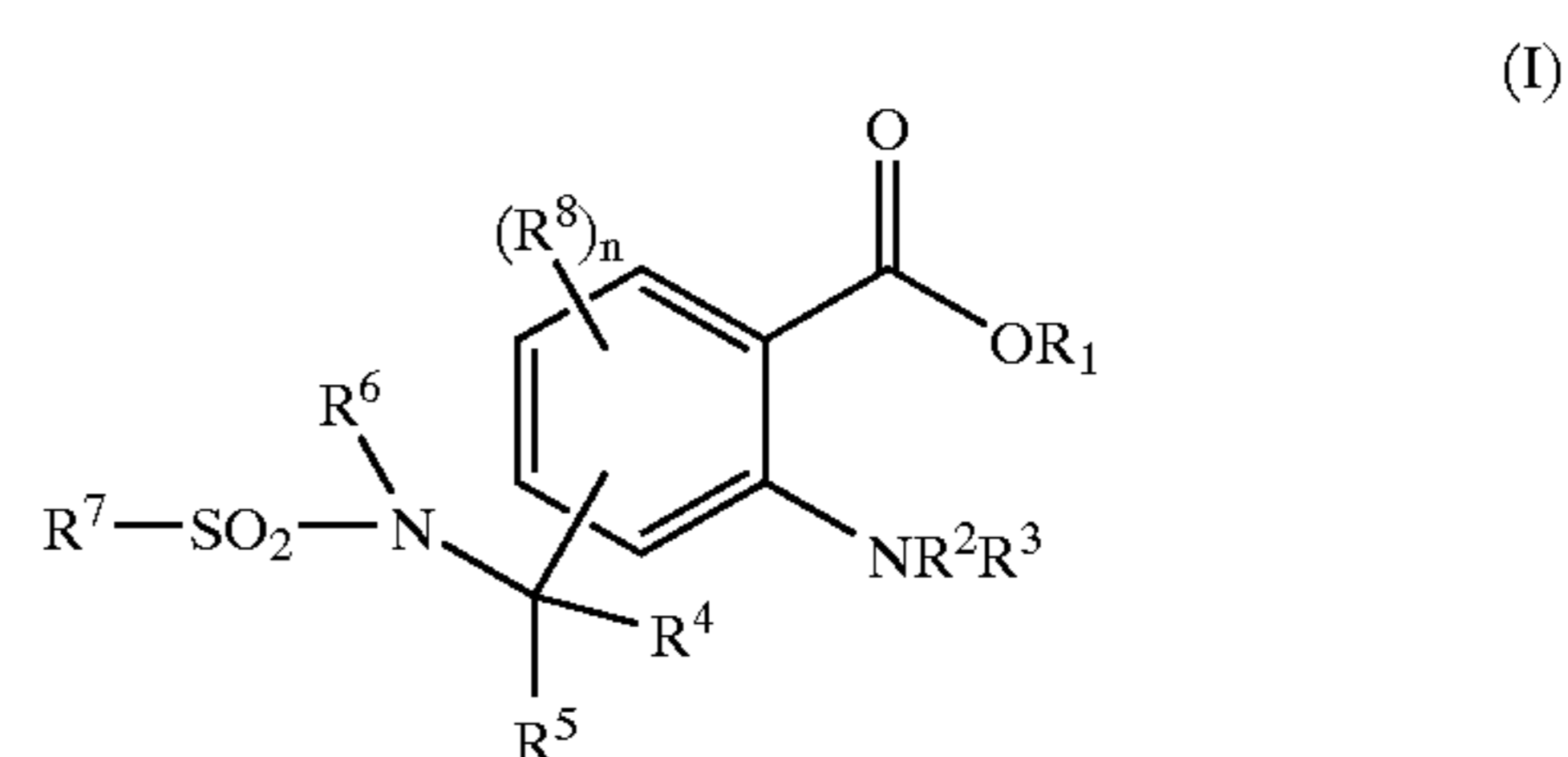
a) reacting a compound of the formulae



with either a reagent that reduces a nitro group to an amine or by catalytic hydrogenation to give a compound of the formula



b) reacting the compound of formula (III) or (IIIa) with a sulfonic acid derivative to give a compound of the formula

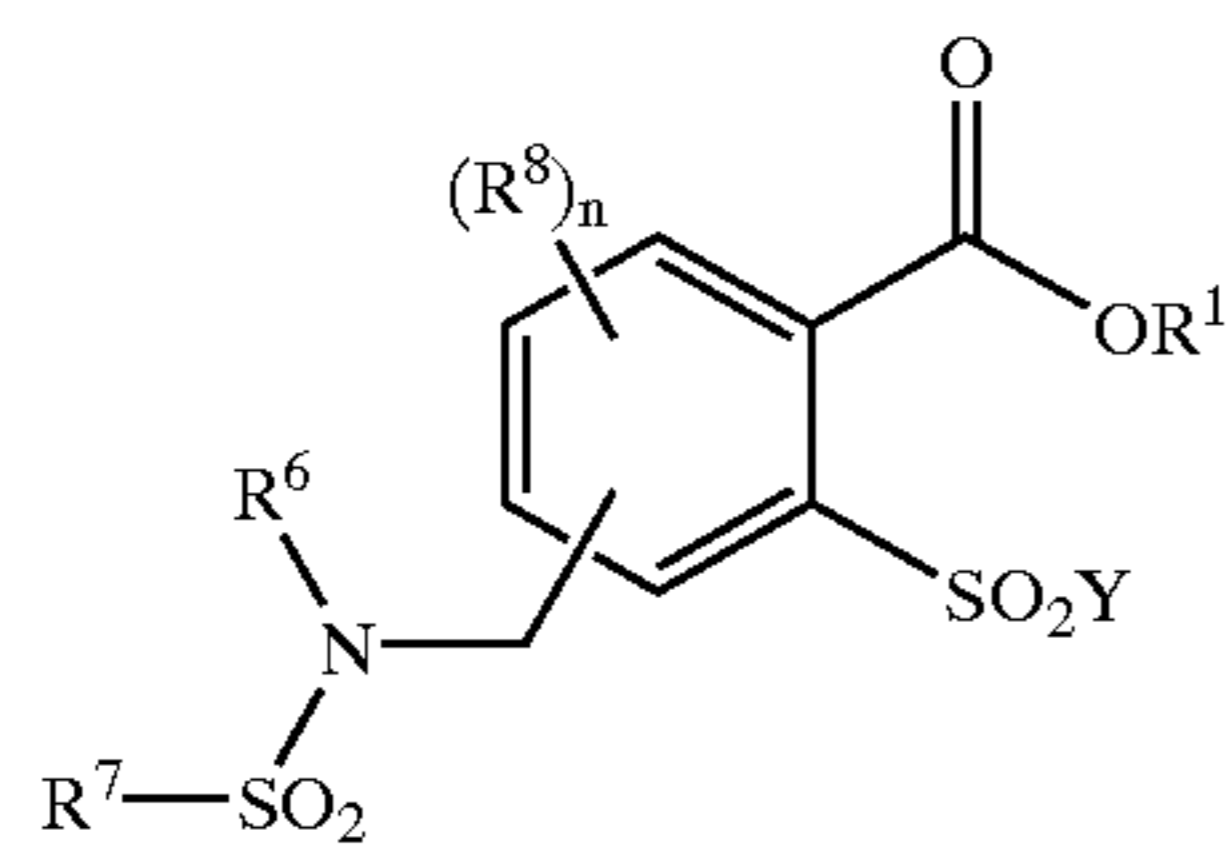


which R¹, R², and R⁶ are hydrogen

c) reacting the compound of formula (I) with an alkylating agent or by reductive amination to give a compound of the formula (I) where R⁶ is an unsubstituted or substituted C₁-C₈-alkyl group

d) first reacting the compound formed in step c) with a diazotizing agent in the presence of an acid followed by an SO₂ source in the presence of a copper catalyst and an acid to give a compound of the formula

35

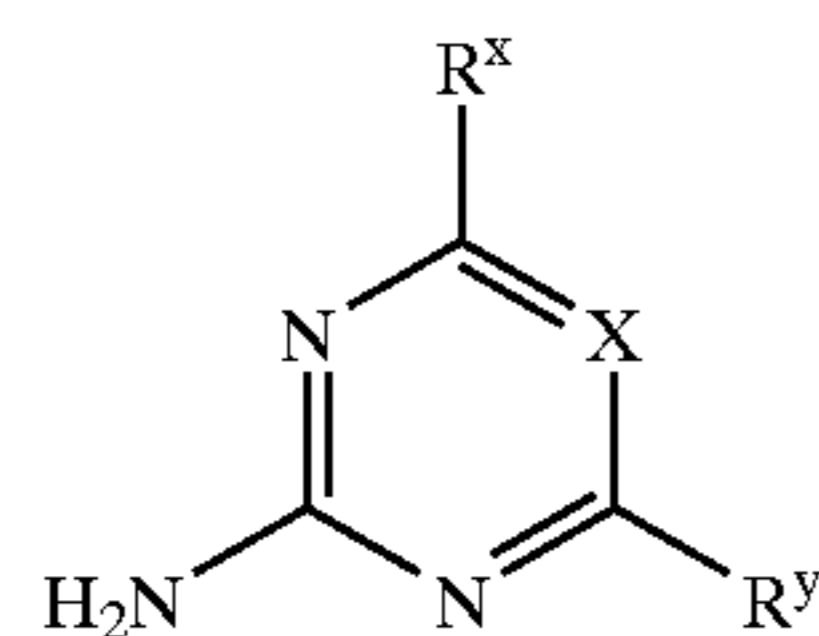


(VII)

where Y is halogen

e) reacting the compound of formula (VII) with either:

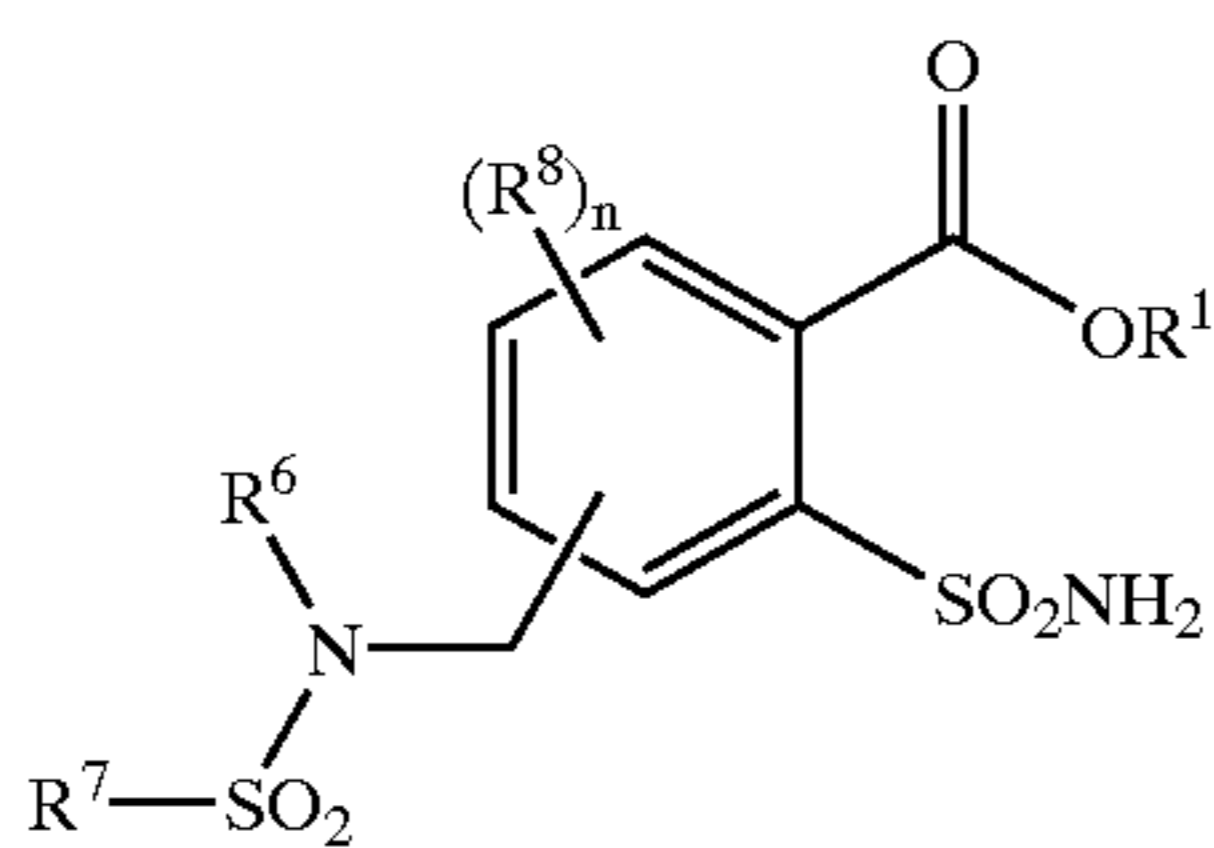
i) amine of the formula (XII) in the presence of MOCN, wherein M is an ammonium ion or an alkali metal ion, to give a compound of the formula (XIII)



(XII)

or

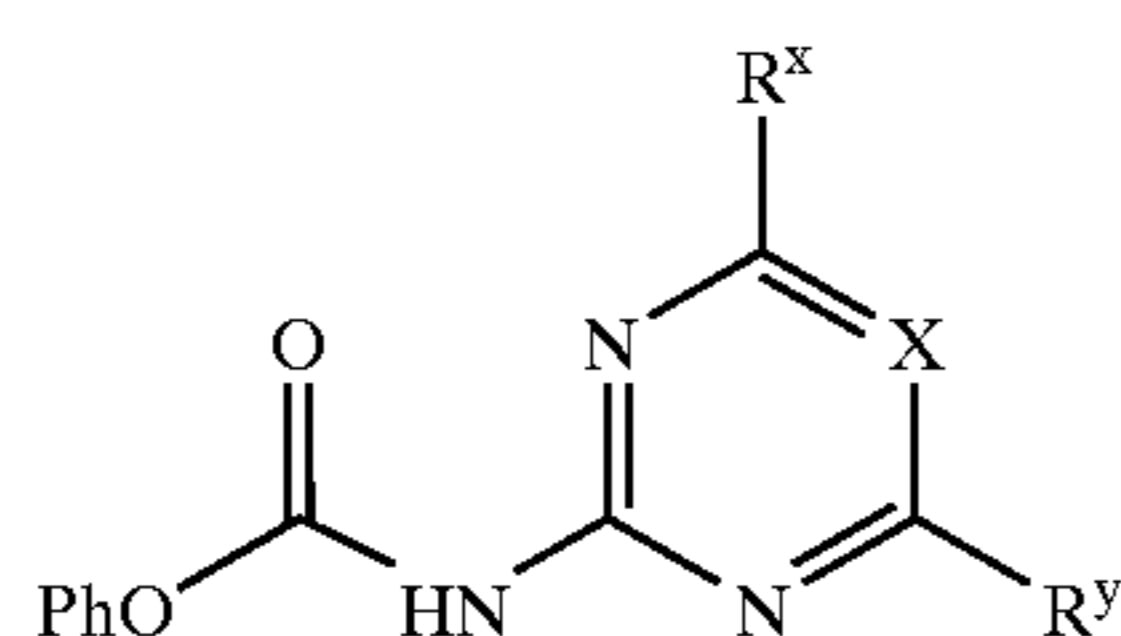
ii) ammonia in the presence of a suitable solvent to give a compound



(VIII)

and subsequently preparing the compound of the formula (XIII) by

a) reacting the compound of the formula (VIII) with a compound of the formula (IX) to give the compound of the formula (XIII)

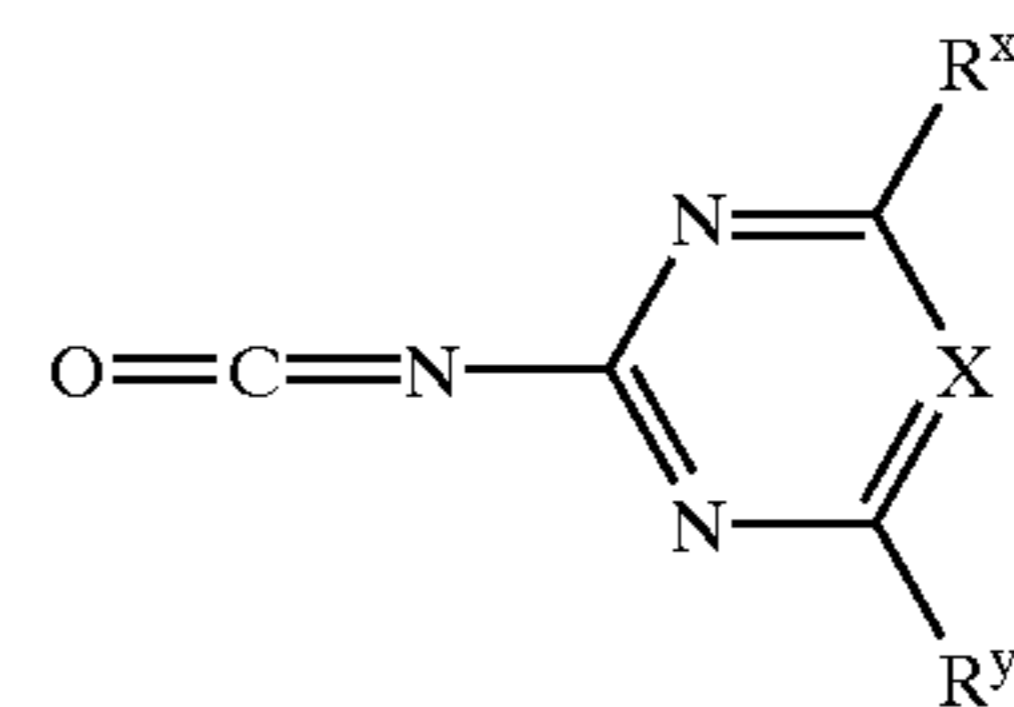


(IX)

or

b) reacting the compound of the formula (VIII) with an isocyanate of the formula (X) to give the compound of the formula (XIII)

36



(X)

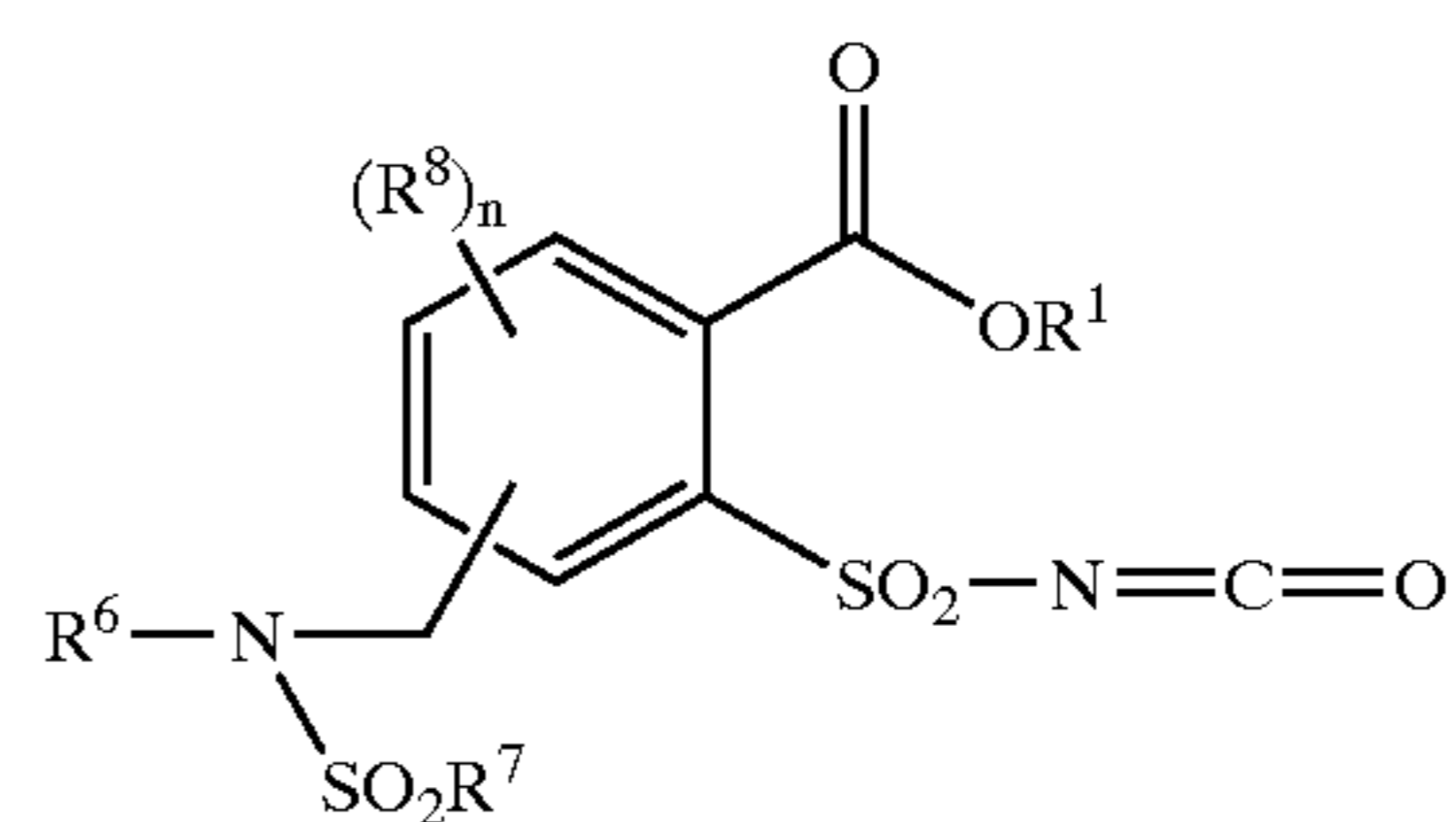
5

10

or

c) reacting the compound of the formula (VIII) with an alkyl isocyanate and phosgene to give a compound of the formula (XI)

15



(XI)

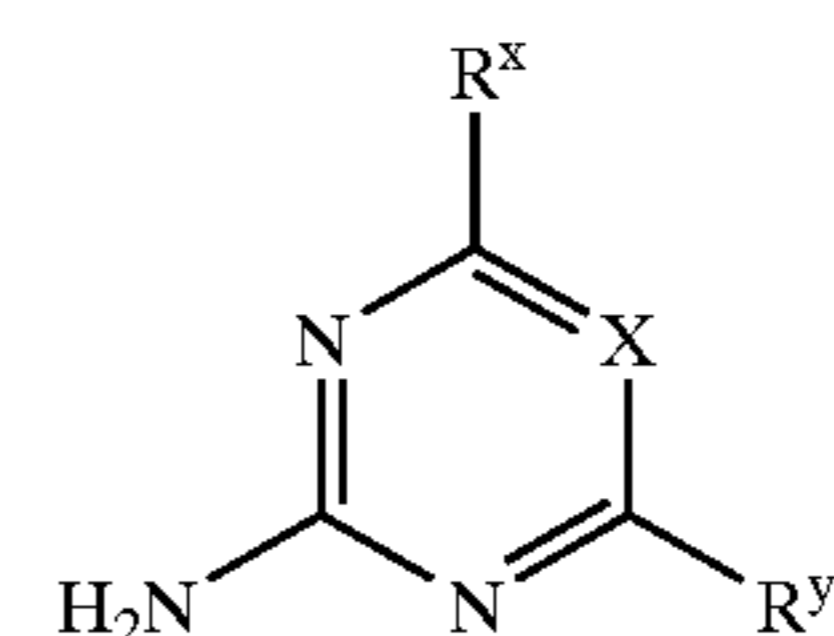
20

25

which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

(XII)

30



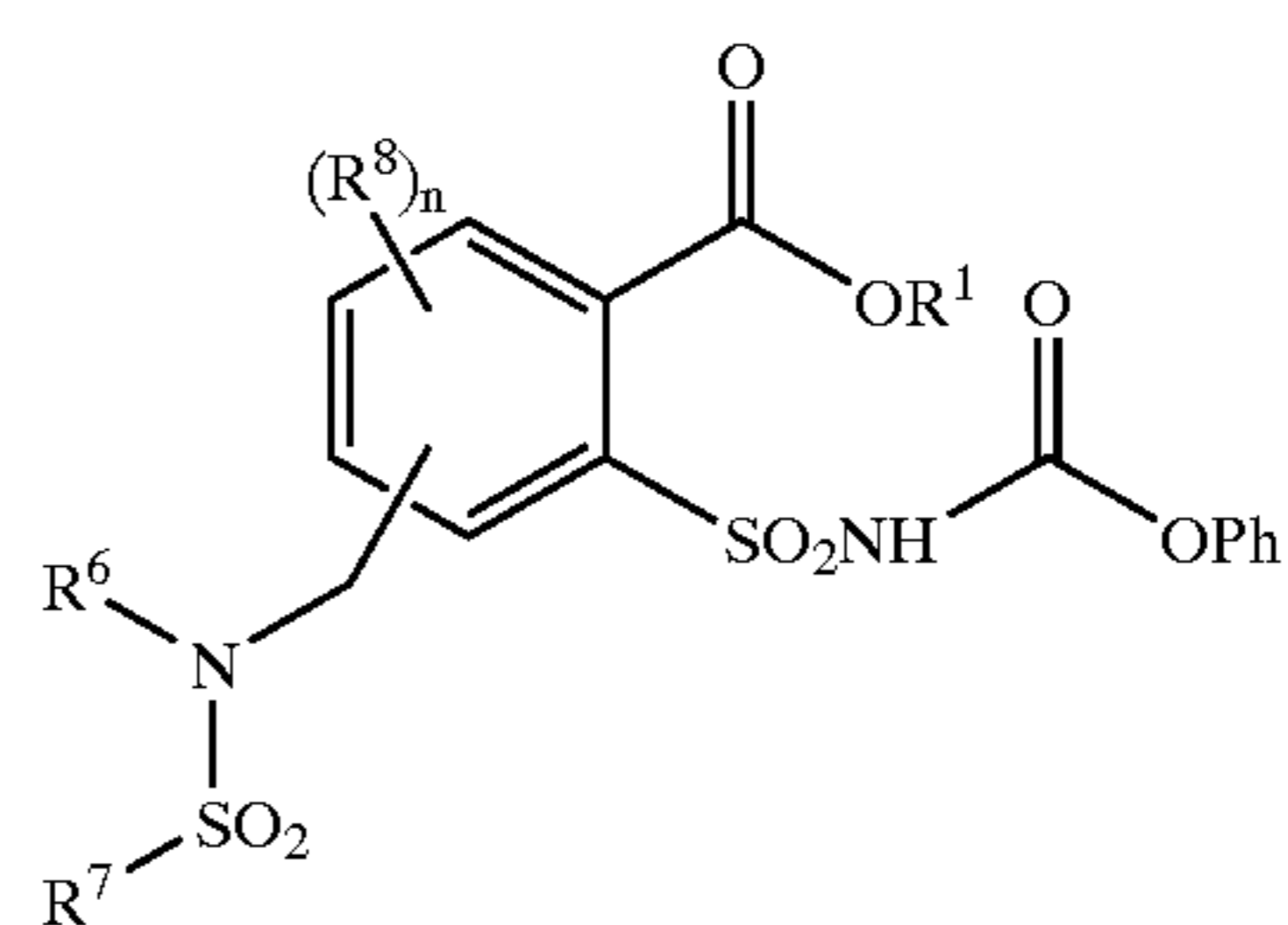
35

or

d) reacting the compound of the formula (VIII) with a carbonic acid derivative R-CO-OPh, in which Ph=unsubstituted or substituted phenyl and R=halogen or unsubstituted or substituted phenoxy to give a compound of the formula (XIV)

40

45



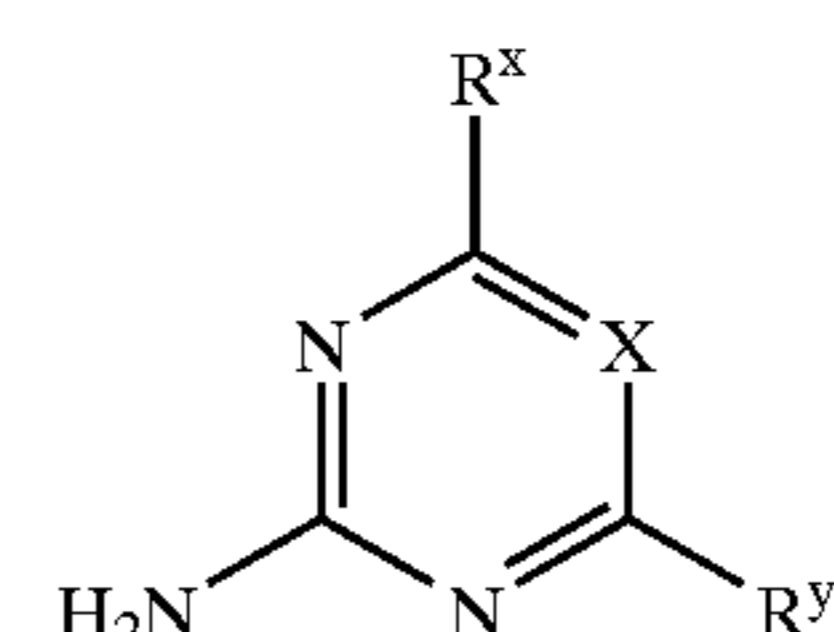
(XIV)

50

which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

(XII)

60



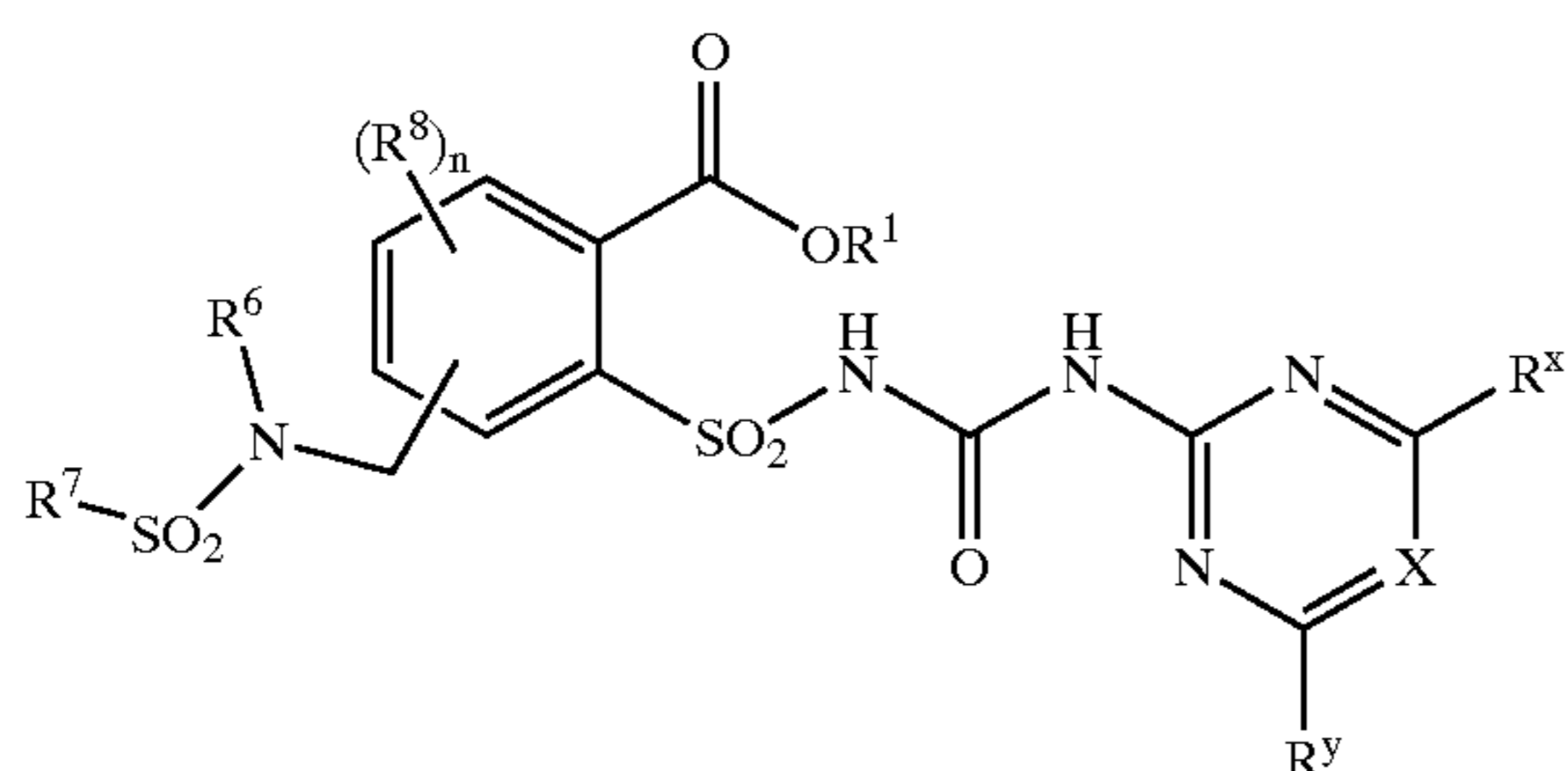
where R¹, R⁷, R⁸, and n in formulae (XI) and (XIV) are as defined in formula (1) R^x, R^y and X in formulae (IX), (X)

37

and (XII) are as defined in formula (XIII) of the present claim and Ph in formulae (IX) and (XIV) is unsubstituted or substituted phenyl, wherein the substituents on the phenyl ring are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₁-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C₁-C₈)alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy.

6. A process of preparing a sulfonyleurea of the formula

(XIII)



in which

R¹ is H, (C₁-C₈)alkyl, (C₃-C₈)alkenyl or (C₃-C₈)alkynyl, where the last 3 radicals are unsubstituted or substituted,

R⁶ is H or (C₁-C₈)alkyl which is unsubstituted or substituted,

R⁷ is (C₁-C₈)alkyl, (C₃-C₈)alkenyl, (C₃-C₈)alkynyl, (C₆-C₁₄)aryl or mono- or di(C₁-C₈)alkylamino which are unsubstituted or substituted, or

R⁶ and R⁷ together form a chain of the formula —(CH₂)_mEm1- which is unsubstituted or substituted, and where m=2, 3 or 4, m¹=0 or 1 and E=CO or SO₂,

R⁸ radicals are identical or different and are (C₁-C₄)alkyl, (C₁-C₄)alkoxy, [(C₁-C₄)alkyl]carbonyl or [(C₁-C₄)alkoxy]carbonyl which are unsubstituted or substituted, or R⁸ is halogen, NO₂ or CN, and

n is 0, 1, 2 or 3,

R^x, R^y independently of one another are a hydrogen atom, halogen, (C₁-C₄)alkyl, (C₁-C₄)alkoxy, (C₁-C₄)alkylthio, where each of the last-mentioned 3 radicals is unsubstituted or substituted by one or more radicals selected from the group consisting of halogen, (C₁-C₄)alkoxy and (C₁-C₄)alkylthio, or are mono- or di[(C₁-C₄)alkyl]amino, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, (C₃-C₆)alkenyloxy or (C₃-C₆)alkynyloxy, and

X is CH or N

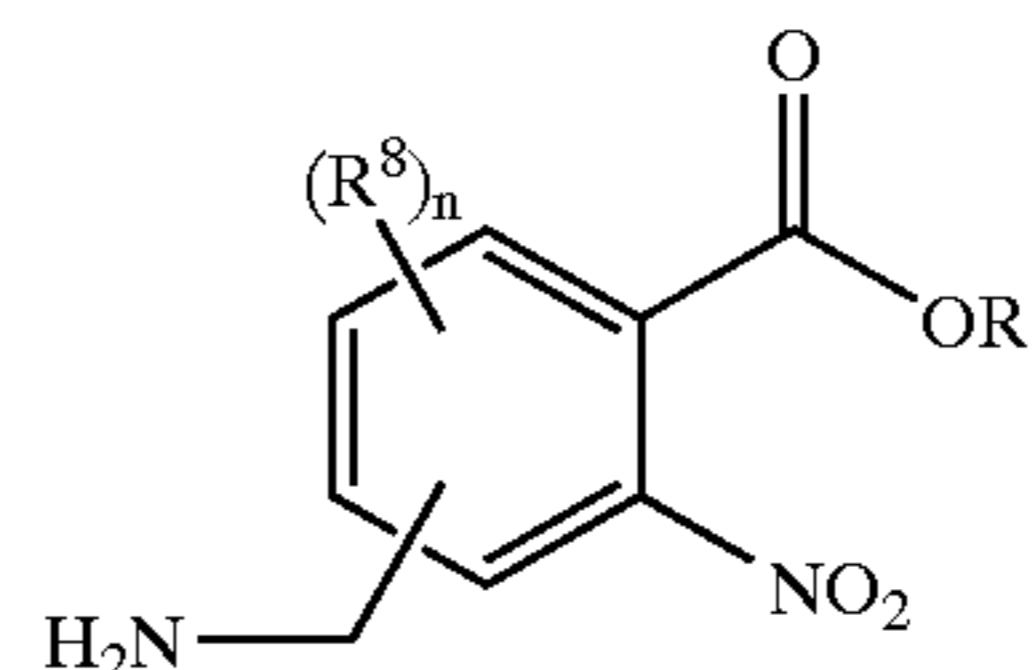
and wherein, unless otherwise identified, the substituents are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₁-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C₁-C₈)

38

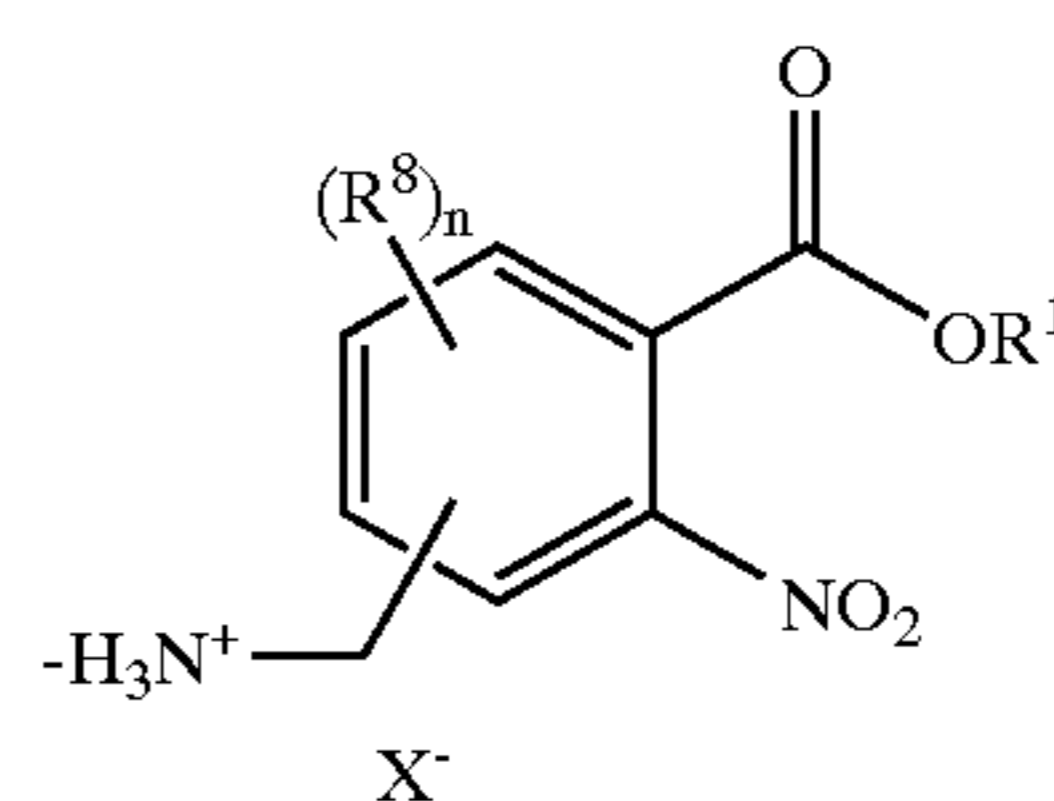
alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy; which comprises:

a) reacting a compound of the formulae

(V)

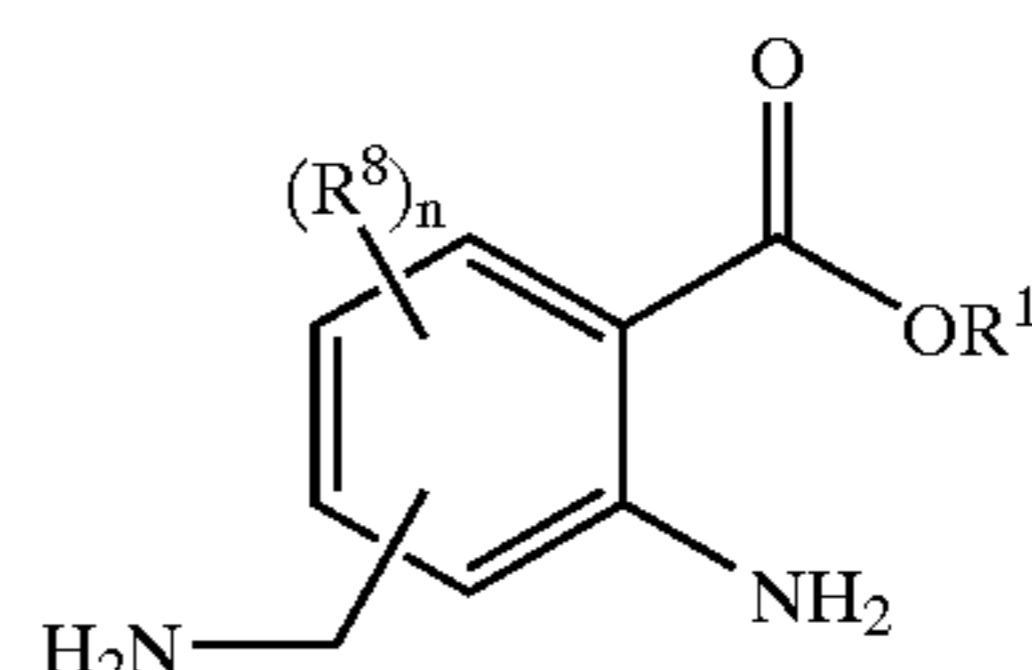


(Va)

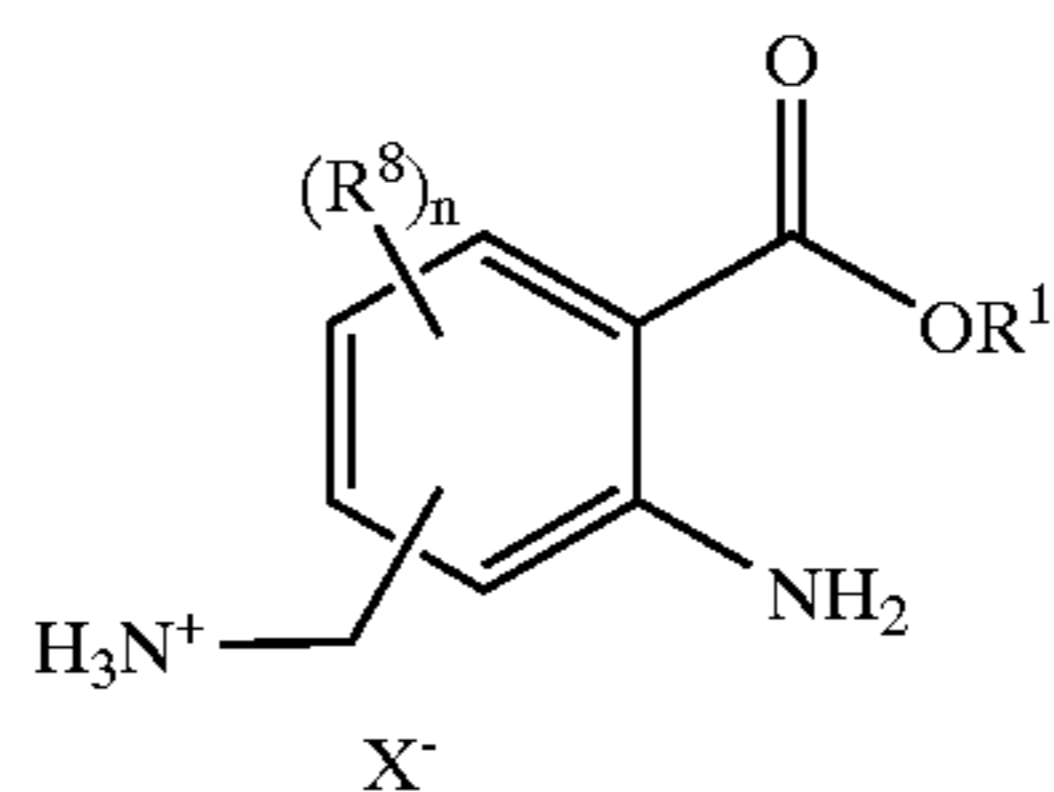


with either a reagent that reduces a nitro group to an amine or by catalytic hydrogenation to give a compound of the formula

(III)

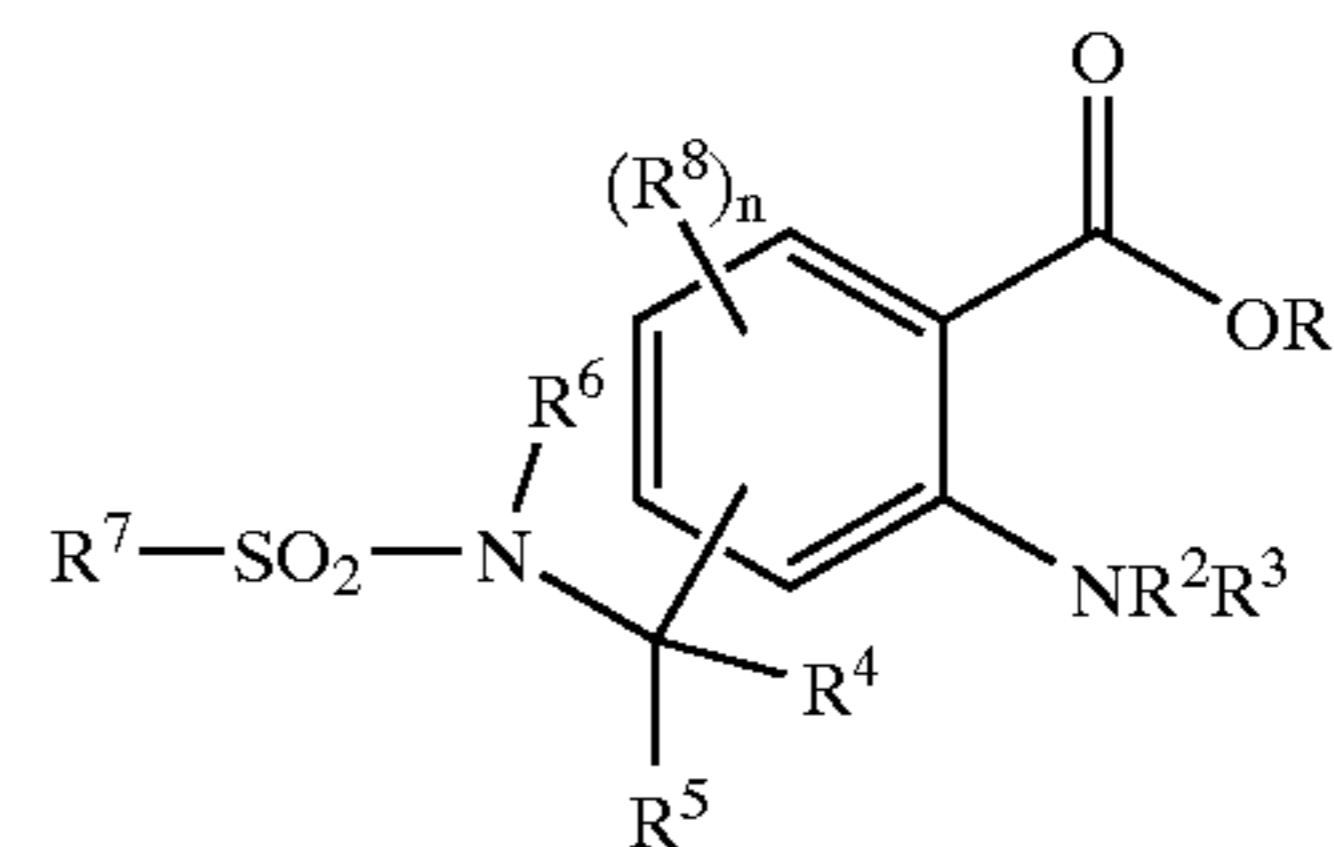


(IIIa)



b) reacting the compound of formula (III) or (IIIa) with a sulfonic acid derivative to give a compound of the formula

(I)

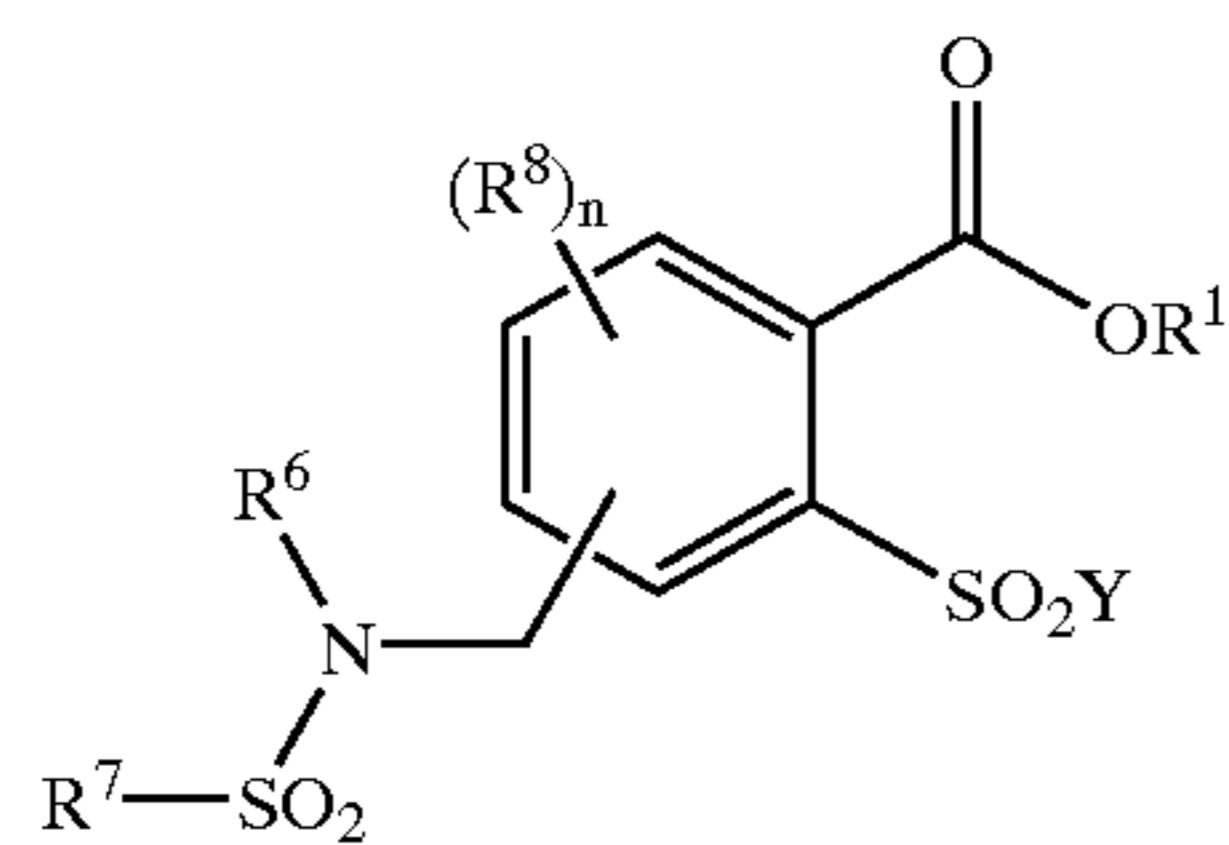


which R¹, R², and R⁶ are hydrogen

c) reacting the compound of formula (I) with an alkylating agent or by reductive amination to give a compound of the formula (I) where R⁶ is an unsubstituted or substituted C₁-C₈-alkyl group

d) first reacting the compound formed in step c) with a diazotizing agent in the presence of an acid followed by an SO₂ source in the presence of a copper catalyst and an acid to give a compound of the formula

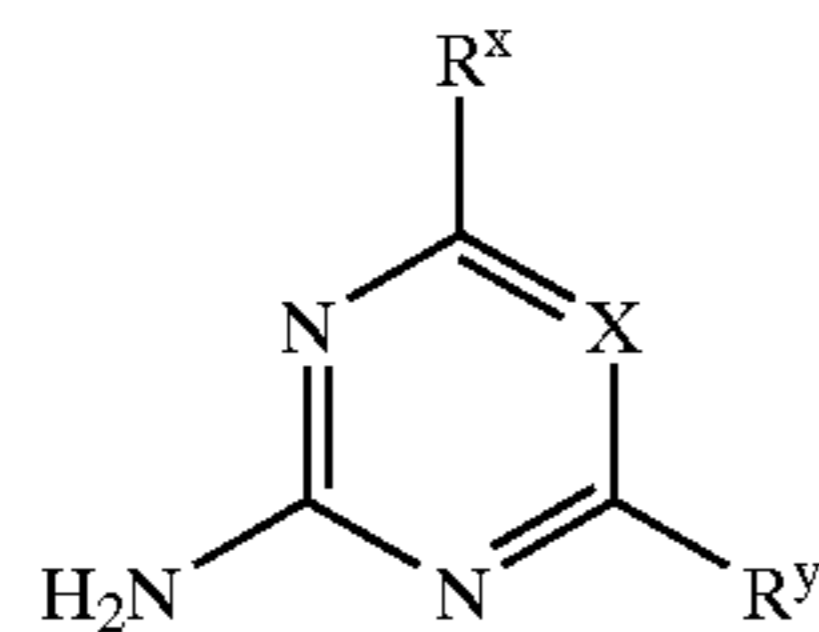
39



where Y is halogen

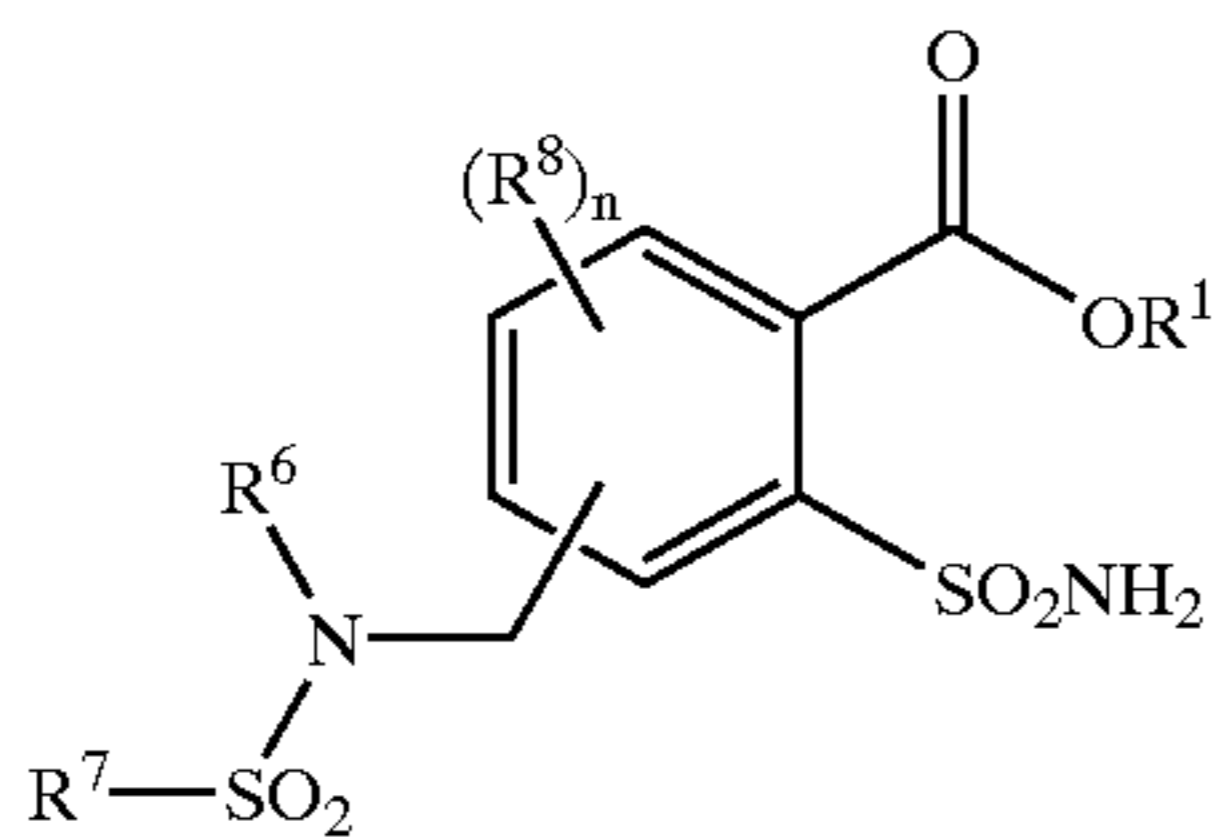
e) reacting the compound of formula (VII) with either:

i) amine of the formula (XII) in the presence of MOCN, wherein M is an ammonium ion or an alkali metal ion, to give a compound of the formula (XIII)



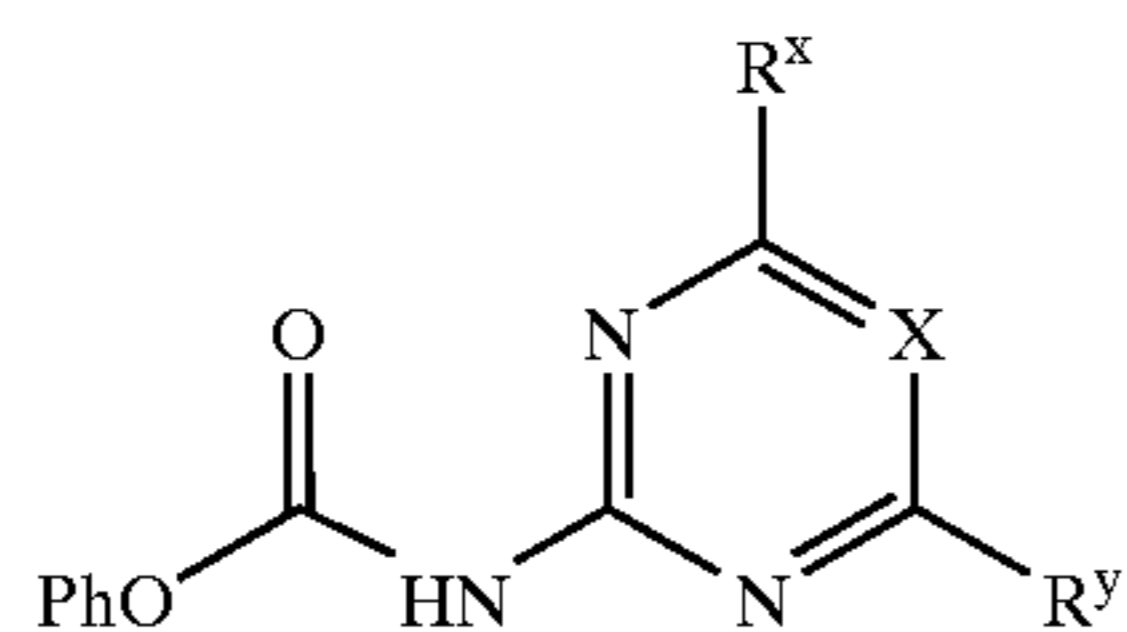
or

ii) ammonia in the presence of a suitable solvent to give a compound



and subsequently preparing the compound of the formula (XIII) by

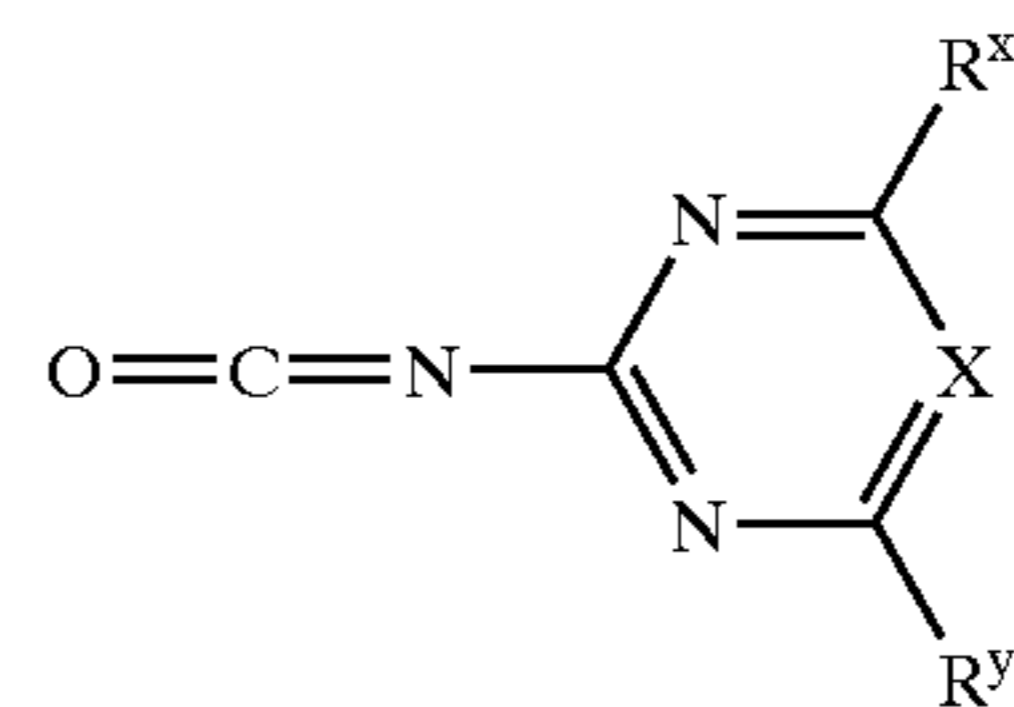
a) reacting the compound of the formula (VIII) with a compound of the formula (IX) to give the compound of the formula (XIII)



or

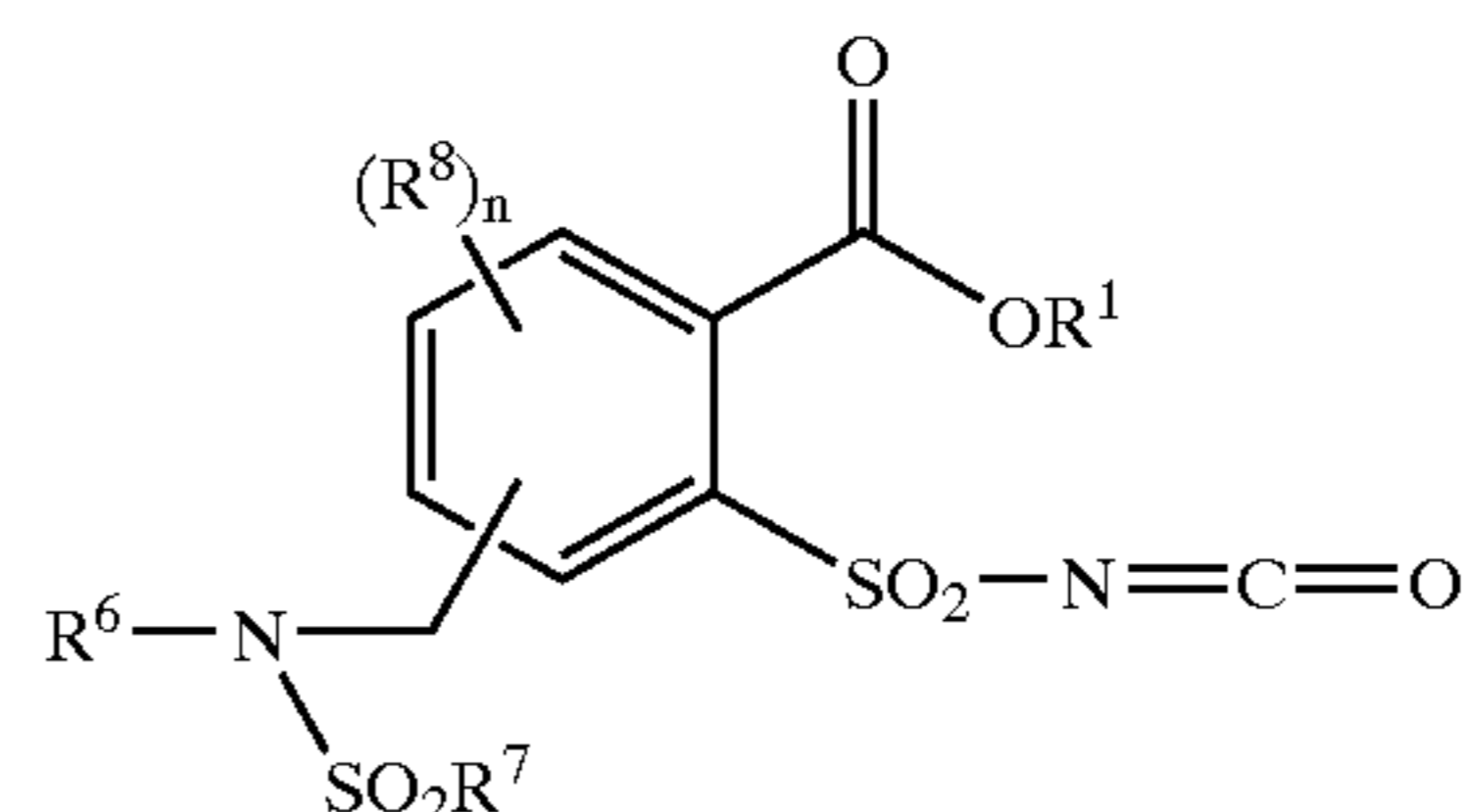
b) reacting the compound of the formula (VIII) with an isocyanate of the formula (X) to give the compound of the formula (XIII)

40

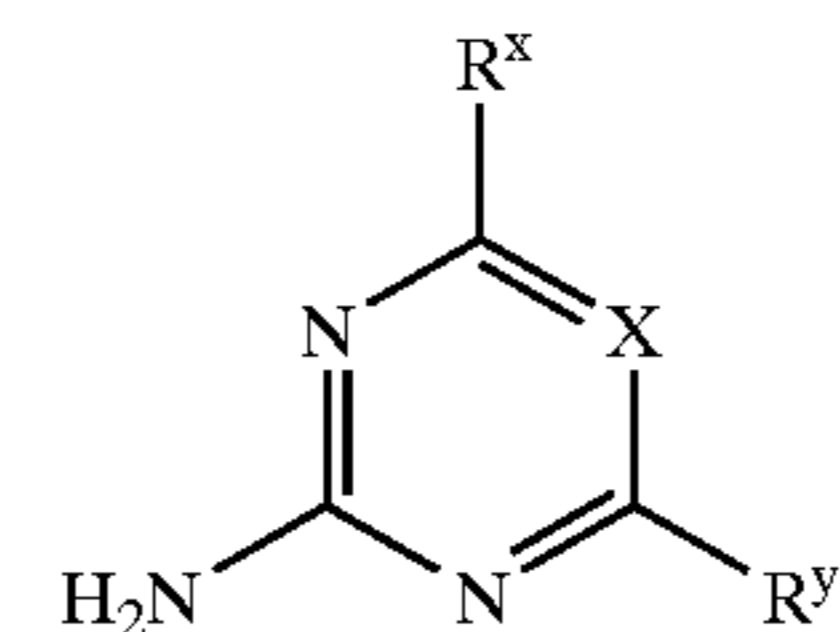


or

c) reacting the compound of the formula (VIII) with an alkyl isocyanate and phosgene to give a compound of the formula (XI)

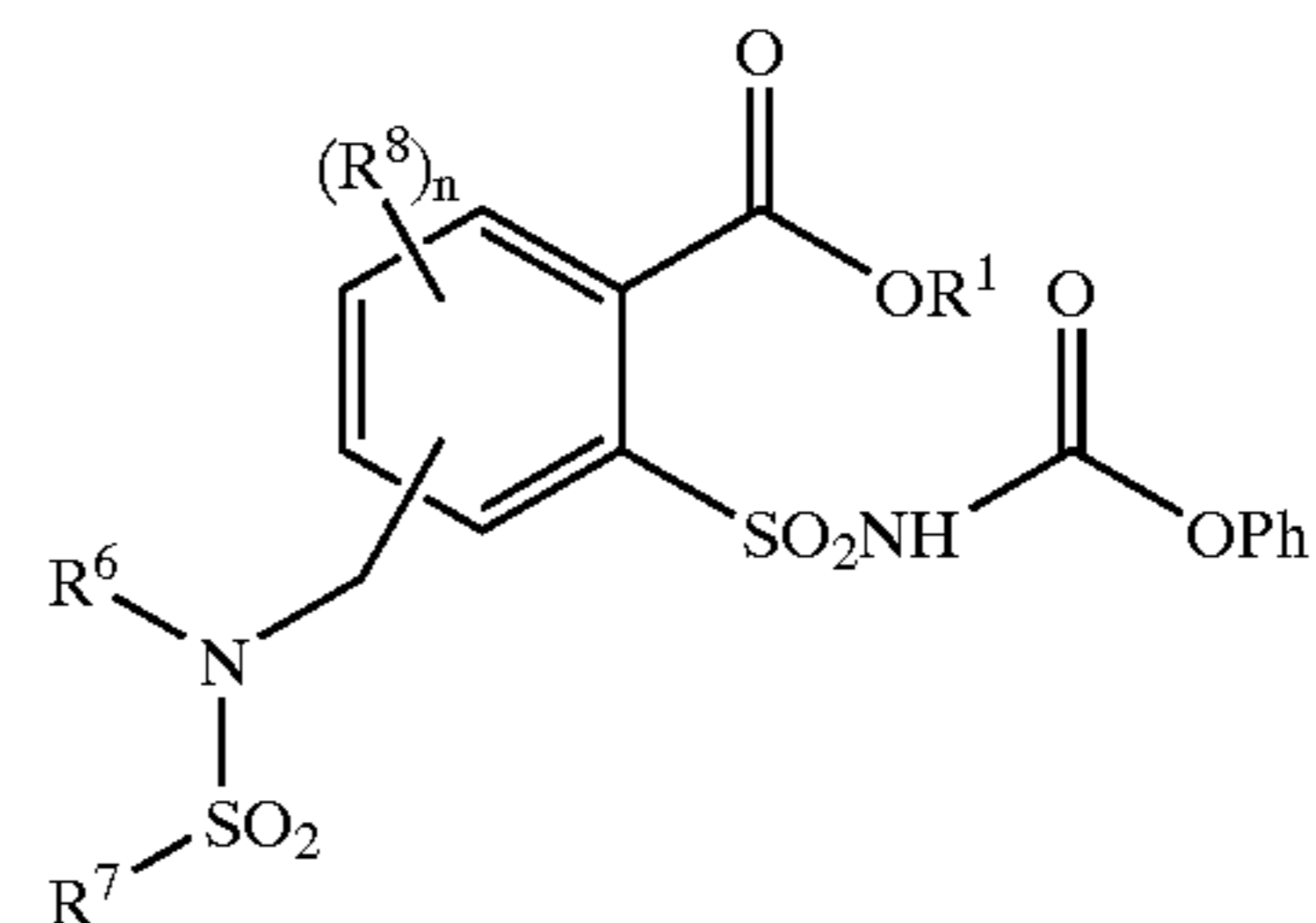


which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)

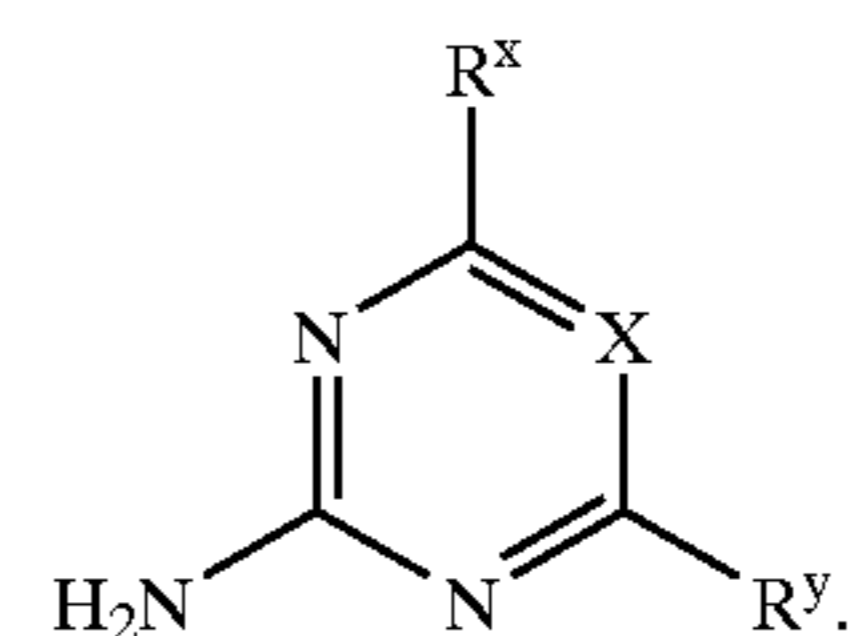


or

d) reacting the compound of the formula (VIII) with a carbonic acid derivative R—CO—OPh, in which Ph=unsubstituted or substituted phenyl and R=halogen or unsubstituted or substituted phenoxy to give a compound of the formula (XIV)

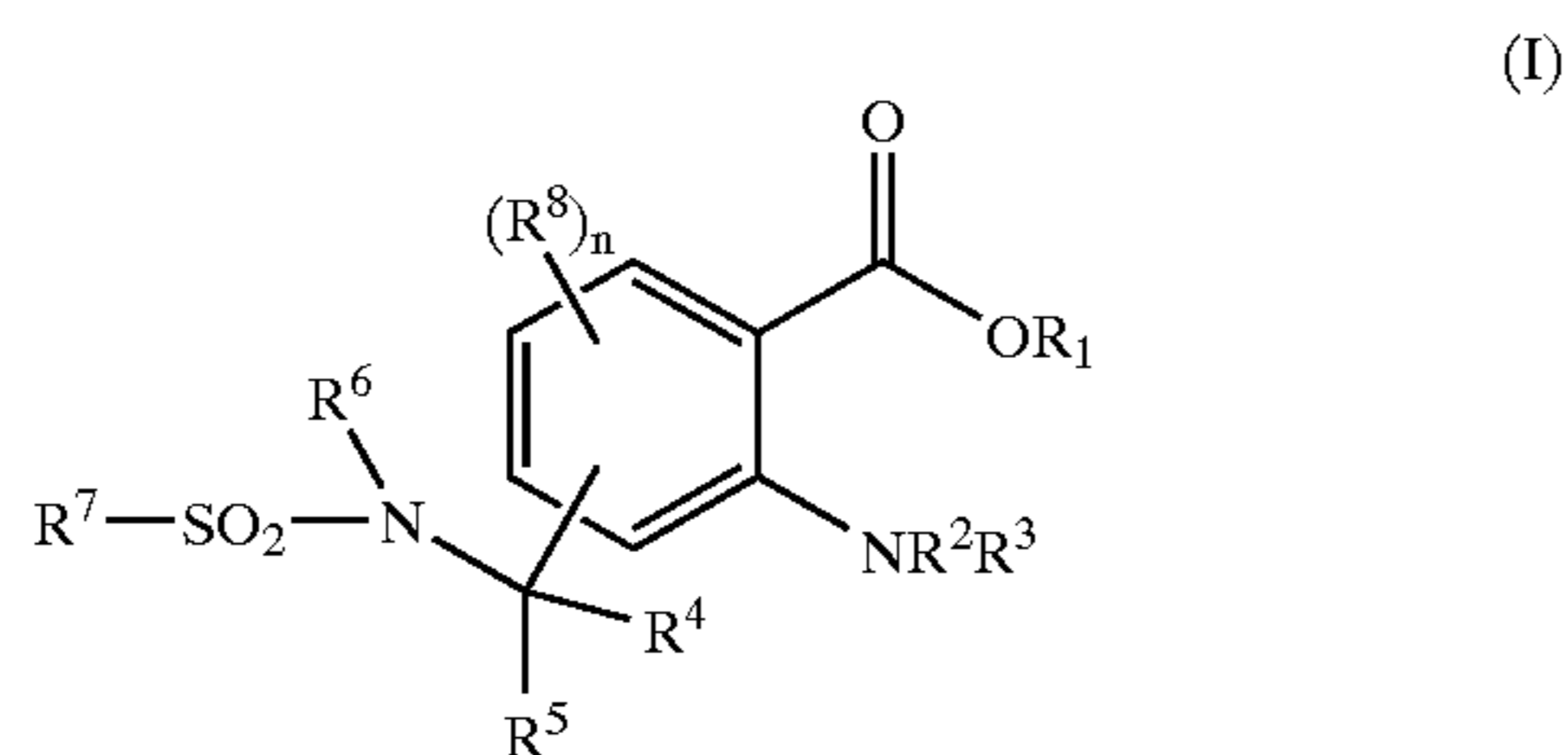


which is subsequently reacted with an amine of the formula (XII) to give the compound of the formula (XIII)



41

7. A process for preparing a compound of the formula



in which

R^1 is H, (C_1-C_8) alkyl, (C_3-C_8) alkenyl or (C_3-C_8) alkynyl, where the last 3 radicals are unsubstituted or substituted,

R^2 , R^3 independently of one another are H or acyl,

R^4 , R^5 are H,

R^6 is H or (C_1-C_8) alkyl which is unsubstituted or substituted,

R^7 is (C_1-C_8) alkyl, (C_3-C_8) alkenyl, (C_3-C_8) alkynyl, (C_6-C_{14}) aryl or mono- or di- (C_1-C_8) alkylamino which are unsubstituted or substituted, or

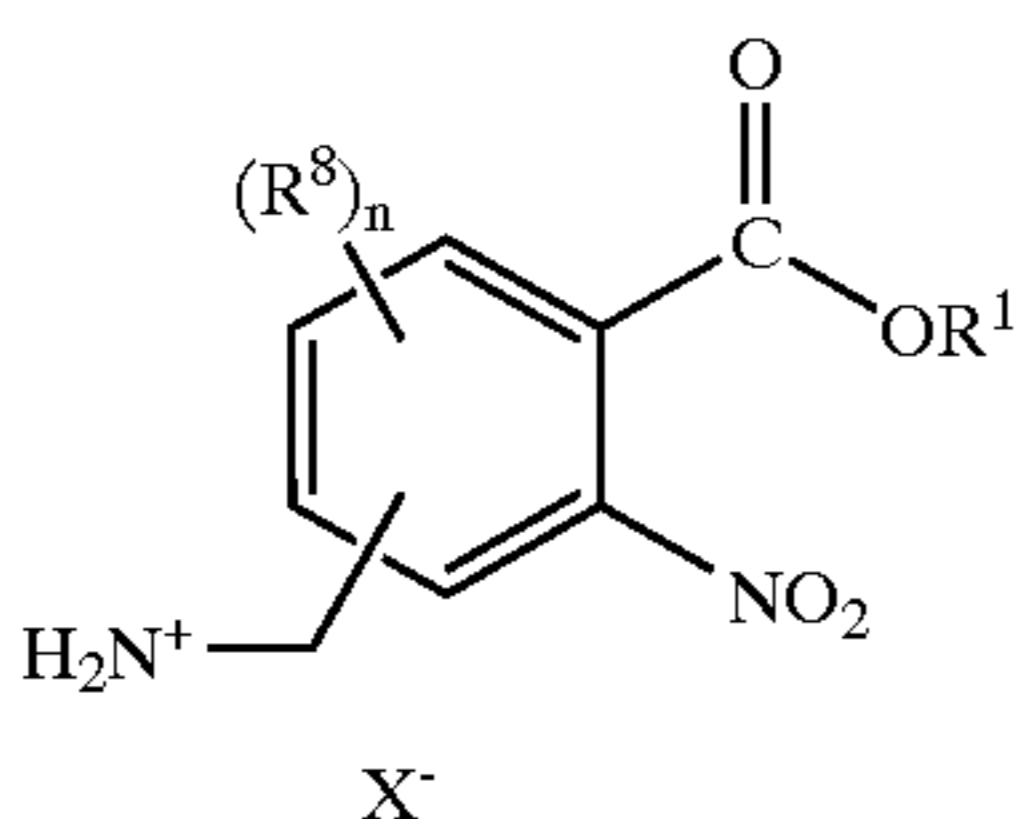
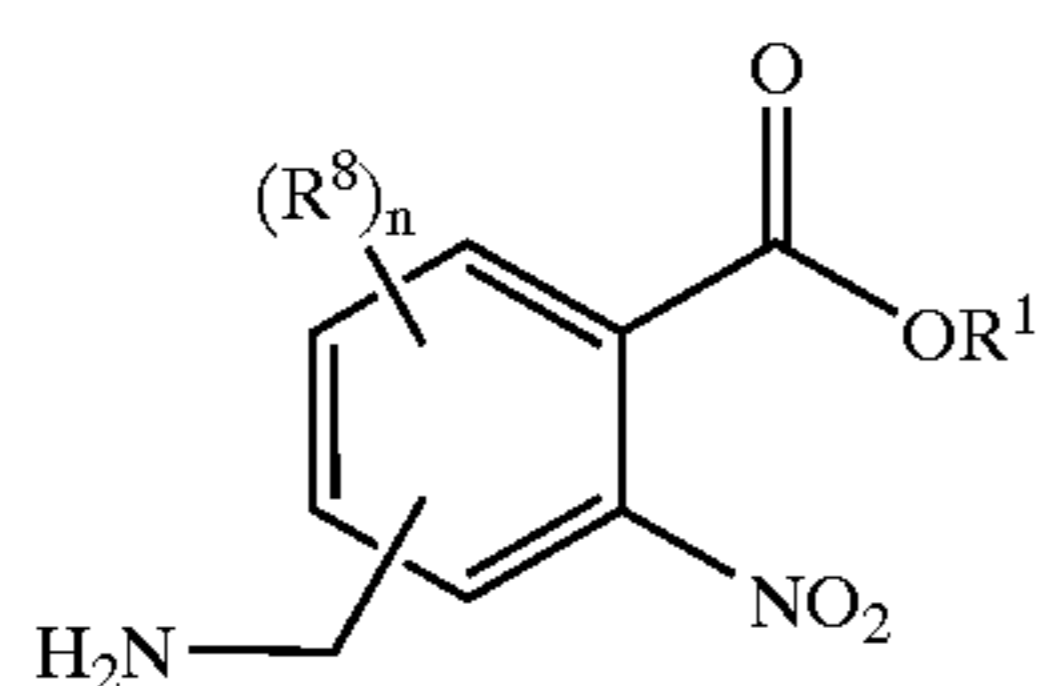
R^6 and R^7 together form a chain of the formula $-(CH_2)_m E_m^1-$ which is unsubstituted or substituted, and where $m=2, 3$ or 4 , $m^1=0$ or 1 and $E=CO$ or SO_2 ,

R^8 radicals are identical or different and are (C_1-C_4) alkyl, (C_1-C_4) alkoxy, $[(C_1-C_4)$ alkyl]carbonyl or $[(C_1-C_4)$ alkoxy]carbonyl which are unsubstituted or substituted, or R^8 is halogen, NO_2 or CN, and

n is 0, 1, 2 or 3,

and wherein, unless otherwise identified, the substituents are halogen, (C_1-C_8) alkoxy, (C_1-C_8) haloalkoxy, (C_1-C_8) alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C_1-C_8) alkoxycarbonyl, (C_1-C_8) alkylcarbonyl, formyl, carbamoyl, mono- and di- (C_1-C_8) alkylaminocarbonyl, acylamino, mono- and di- (C_1-C_8) alkylamino, (C_1-C_8) alkylsulfinyl, (C_1-C_8) haloalkylsulfinyl, (C_1-C_8) alkylsulfonyl, halo- (C_1-C_8) alkylsulfonyl, (C_1-C_8) cycloalkyl, (C_1-C_8) cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C_1-C_8) alkyl and halo- (C_1-C_8) alkyl, (C_1-C_8) alkenyl, (C_1-C_8) alkynyl, (C_1-C_8) alkenyloxy or (C_1-C_8) alkynyloxy, which comprises

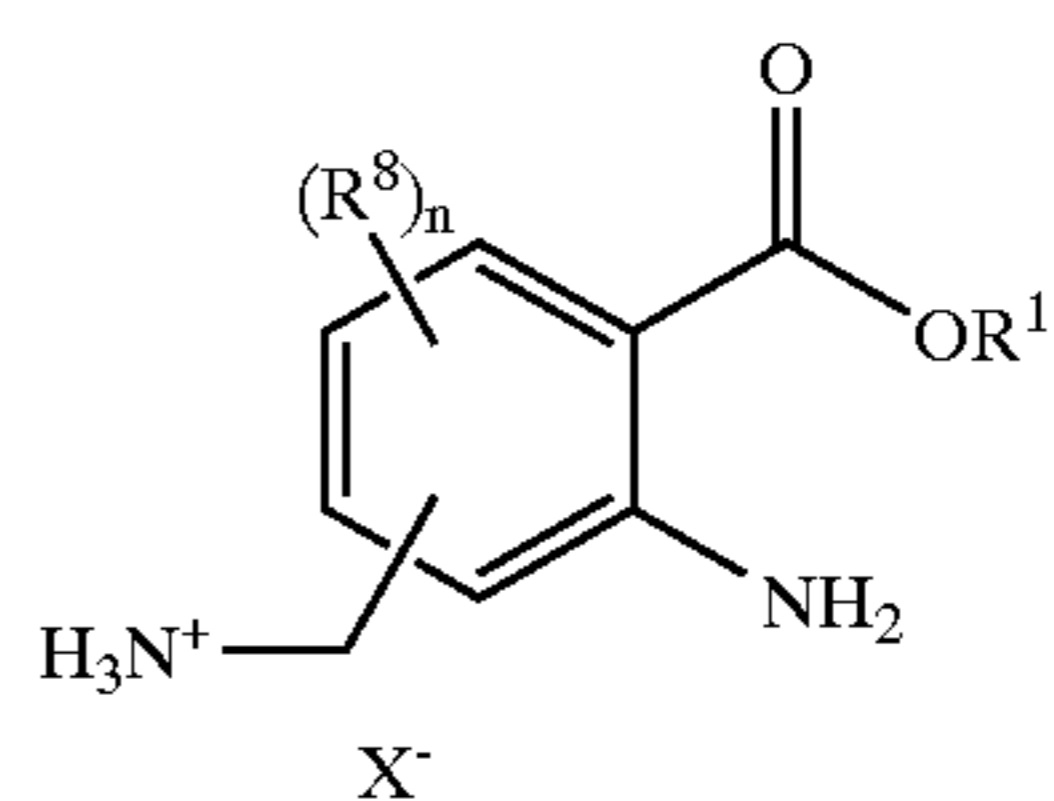
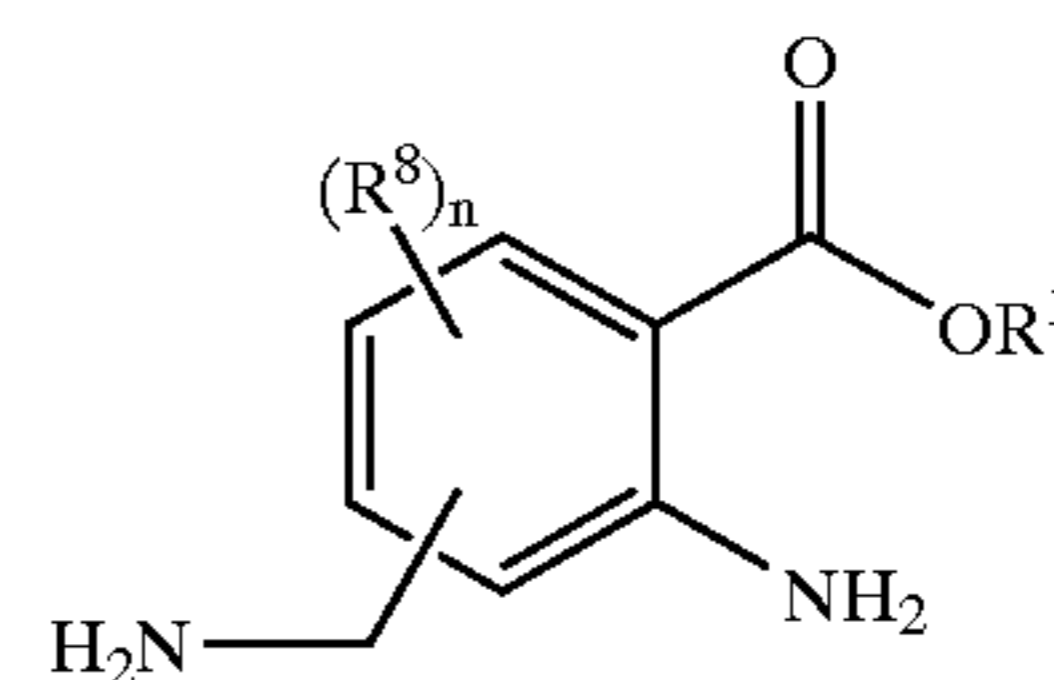
a) reacting a compound of the formulae



with either a reagent that reduces a nitro group to an amine or by catalytic hydrogenation to give a compound of the formula

42

(III)

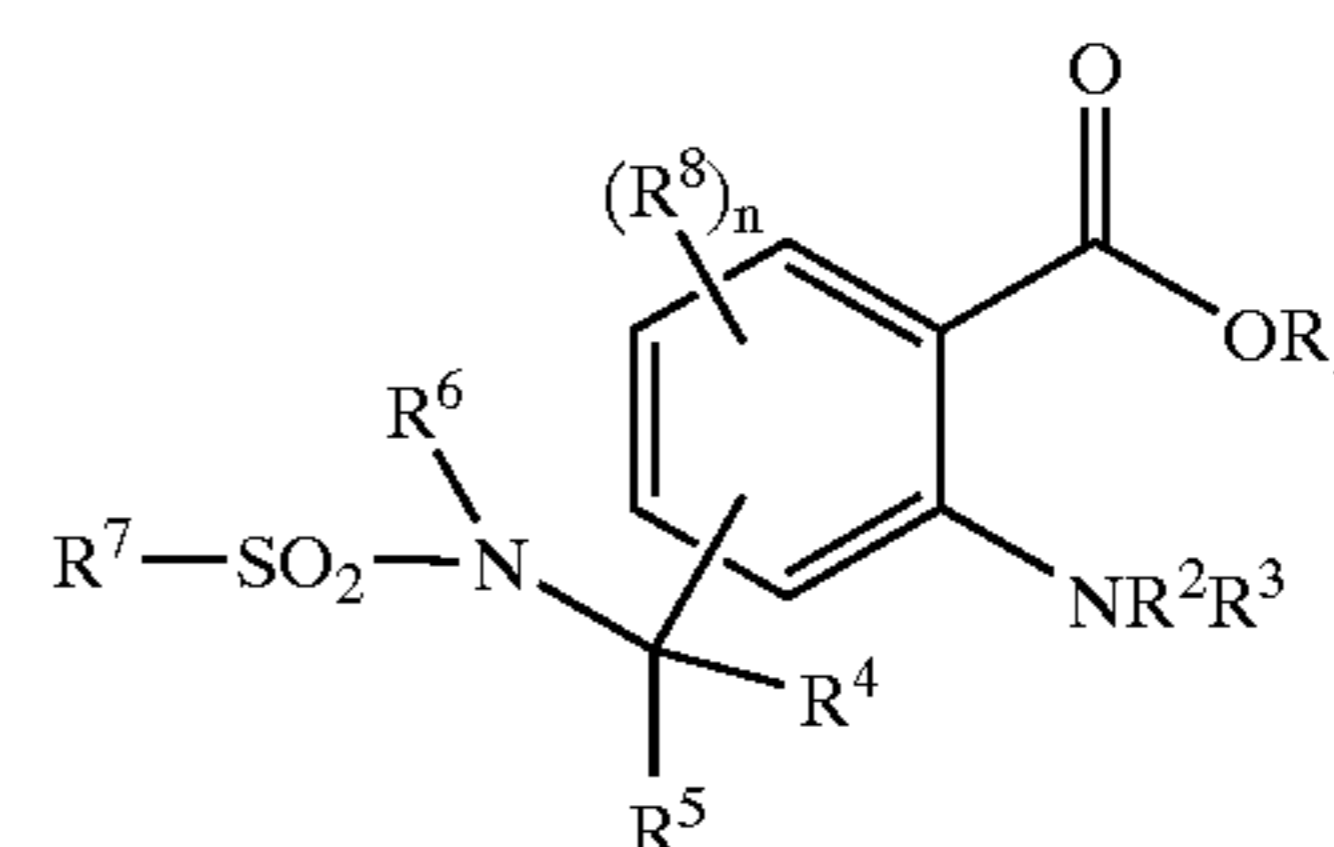


and

b) reacting the compound of formula (III) or (IIIa) with a sulfonic acid derivative to give a compound of the formula (I), wherein R^1 , R^2 , and R^6 are hydrogen, and

c) optionally reacting a compound formed in b) with an alkylating agent or by reductive amination to give a compound of the formula (I) where R^6 an unsubstituted or substituted C_1-C_8 -alkyl group.

8. A process for preparing a compound of the formula



in which

R^1 is H, (C_1-C_8) alkyl, (C_3-C_8) alkenyl or (C_3-C_8) alkynyl, where the last 3 radicals are unsubstituted or substituted,

R^2 , R^3 independently of one another are H or acyl,

R^4 , R^5 are H,

R^6 is H or (C_1-C_8) alkyl which is unsubstituted or substituted,

R^7 is (C_1-C_8) alkyl, (C_3-C_8) alkenyl, (C_3-C_8) alkynyl, (C_6-C_{14}) aryl or mono- or di- (C_1-C_8) alkylamino which are unsubstituted or substituted, or

R^6 and R^7 together form a chain of the formula $-(CH_2)_m E_m^1-$ which is unsubstituted or substituted, and where $m=2, 3$ or 4 , $m^1=0$ or 1 and $E=CO$ or SO_2 ,

R^8 radicals are identical or different and are (C_1-C_4) alkyl, (C_1-C_4) alkoxy, $[(C_1-C_4)$ alkyl]carbonyl or $[(C_1-C_4)$ alkoxy]carbonyl which are unsubstituted or substituted, or R^8 is halogen, NO_2 or CN, and

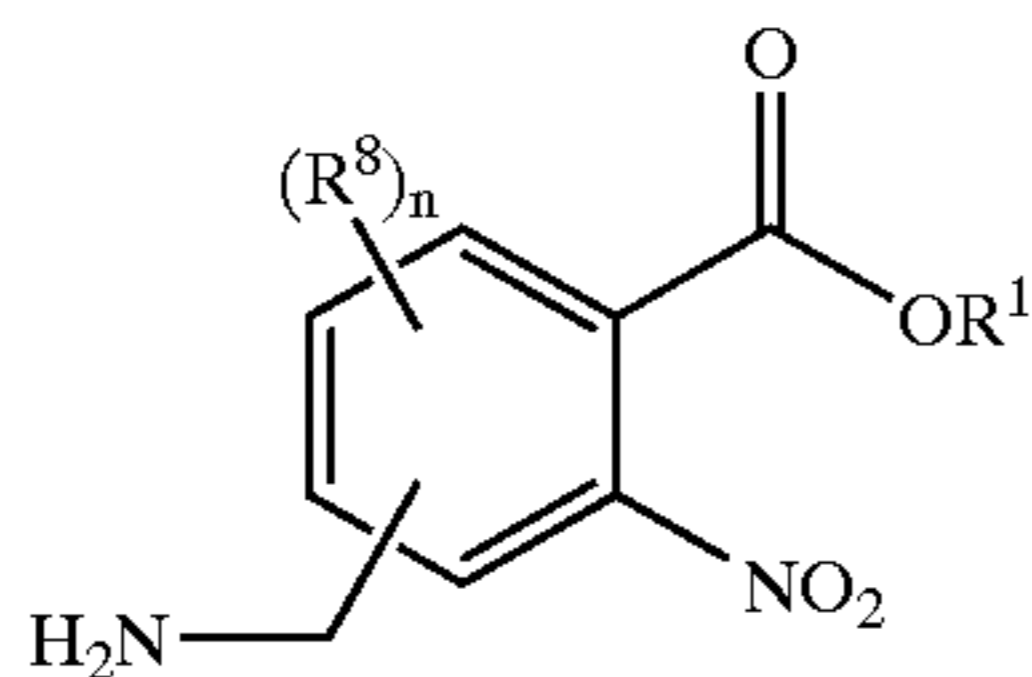
n is 0, 1, 2 or 3,

and wherein, unless otherwise identified, the substituents are halogen, (C_1-C_8) alkoxy, (C_1-C_8) haloalkoxy, (C_1-C_8) alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C_1-C_8) alkoxycarbonyl, (C_1-C_8) alkylcarbonyl, formyl, carbamoyl, mono- and di- (C_1-C_8) alkylaminocarbonyl, acylamino, mono- and di- (C_1-C_8) alkylamino, (C_1-C_8) alkylsulfinyl, (C_1-C_8) haloalkylsulfinyl, (C_1-C_8) alkylsulfonyl, halo- (C_1-C_8) alkylsulfonyl, (C_1-C_8) cycloalkyl, (C_1-C_8) cycloalkenyl, wherein the cycloalkyl

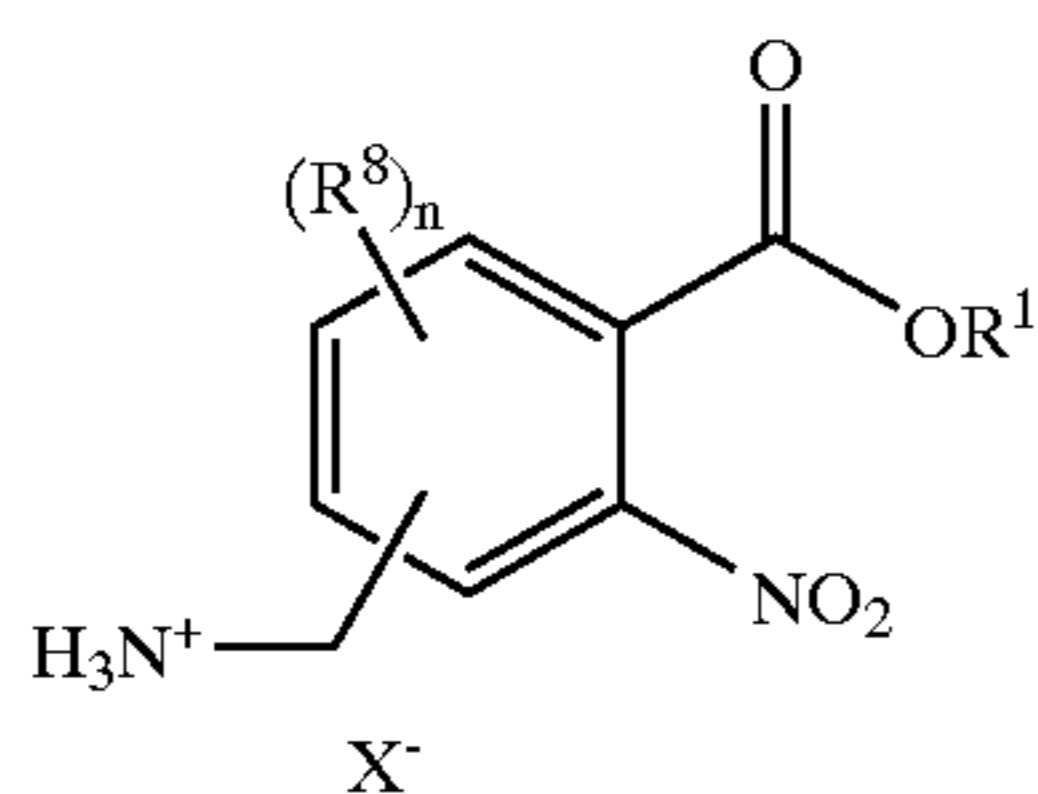
43

and cycloalkyl radicals are optionally substituted by (C₁-C₈) alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy which comprises

a) reacting a compound of the formulae

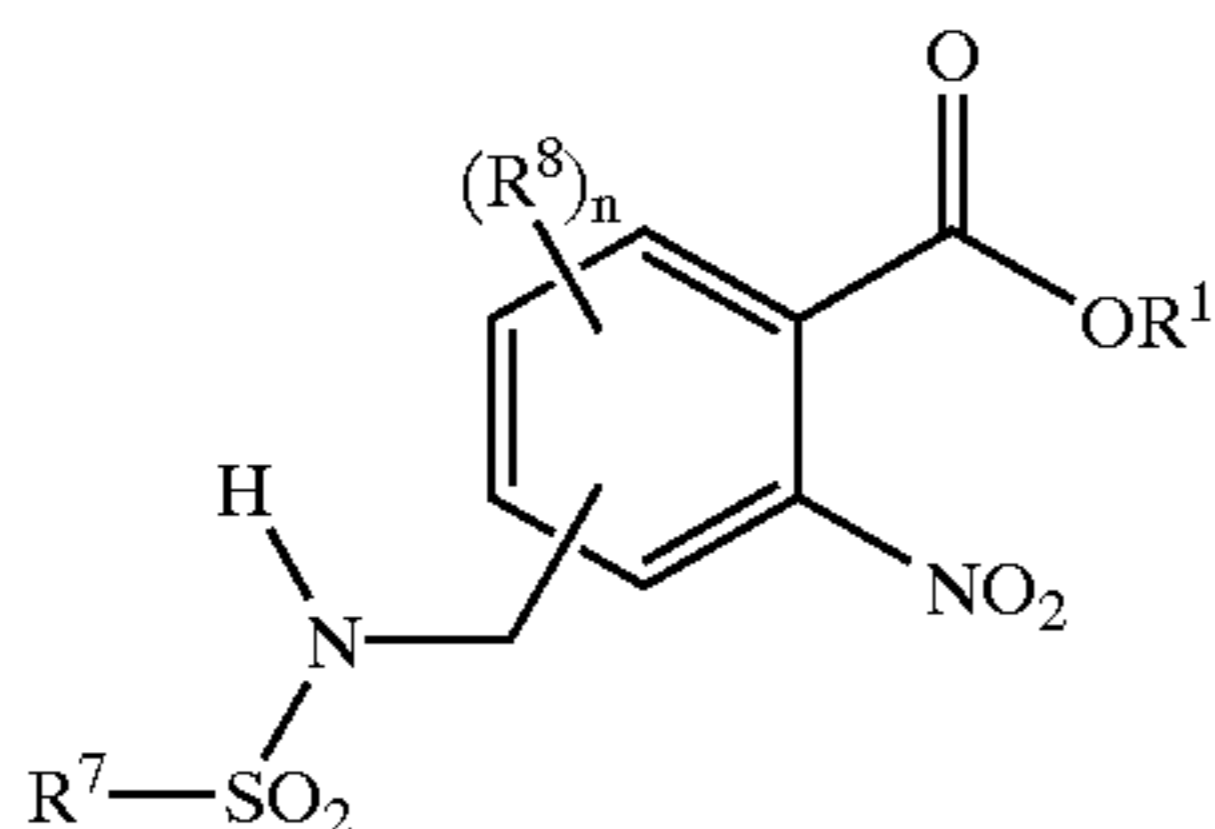


(V)



(Va)

with a sulfonic acid derivative to give a compound of the formula

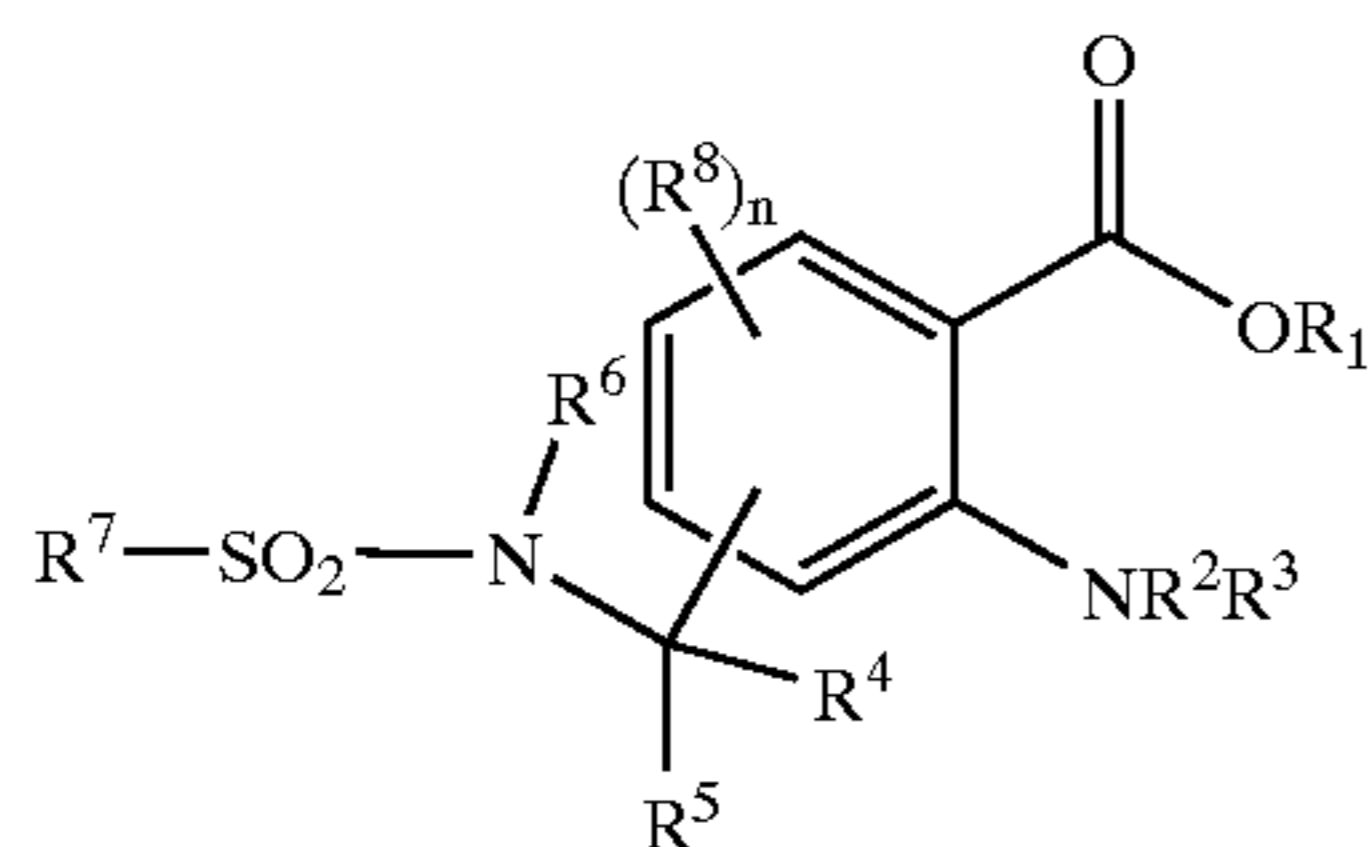


(VI)

and reacting the compound of formula (VI) with an agent which reduces a nitro compound to an amine or by catalytic hydrogenation to give a compound of formula (I), wherein R¹, R², and R⁶ are hydrogen, and

b) optionally reacting a compound of formula (I) formed in a) with an alkylating agent or by reductive amination to give a compound of the formula (I) where R⁶ an unsubstituted or substituted C₁-C₈-alkyl group.

9. A process for preparing a compound of the formula



(I)

in which

R¹ is H, (C₁-C₈)alkyl, (C₃-C₈)alkenyl or (C₃-C₈)alkynyl, where the last 3 radicals are unsubstituted or substituted,

44

R², R³ independently of one another are H or acyl,

R⁴, R⁵ are H,

R⁶ is H or (C₁-C₈)alkyl which is unsubstituted or substituted,

5

R⁷ is (C₁-C₈)alkyl, (C₃-C₈)alkenyl, (C₃-C₈)alkynyl, (C₆-C₁₄)aryl or mono- or di(C₁-C₈)alkylamino which are unsubstituted or substituted, or

(V)

10

R⁶ and R⁷ together form a chain of the formula -(CH₂)^mEm¹- which is unsubstituted or substituted, and where m=2, 3 or 4, m¹=0 or 1 and E=CO or SO₂,

R⁸ radicals are identical or different and are (C₁-C₄)alkyl, (C₁-C₄)alkoxy, [(C₁-C₄)alkyl]carbonyl or [(C₁-C₄)alkoxy]carbonyl which are unsubstituted or substituted, or R⁸ is halogen, NO₂ or CN, and

15

n is 0, 1, 2 or 3,

and wherein, unless otherwise identified, the substituents are halogen, (C₁-C₈)alkoxy, (C₁-C₈)haloalkoxy, (C₁-C₈)alkylthio, hydroxyl, amino, nitro, carboxyl, cyano, azido (C₁-C₈)alkoxycarbonyl, (C₁-C₈)alkylcarbonyl, formyl, carbamoyl, mono- and di-(C₁-C₈)alkylaminocarbonyl, acylamino, mono- and di-(C₁-C₈)alkylamino, (C₁-C₈)alkylsulfinyl, (C₁-C₈)haloalkylsulfinyl, (C₁-C₈)alkylsulfonyl, halo(C₁-C₈)alkylsulfonyl, (C₁-C₈)cycloalkyl, (C₁-C₈)cycloalkenyl, wherein the cycloalkyl and cycloalkyl radicals are optionally substituted by (C₁-C₈)alkyl and halo(C₁-C₈)alkyl, (C₁-C₈)alkenyl, (C₁-C₈)alkynyl, (C₁-C₈)alkenyloxy or (C₁-C₈)alkynyloxy,

20

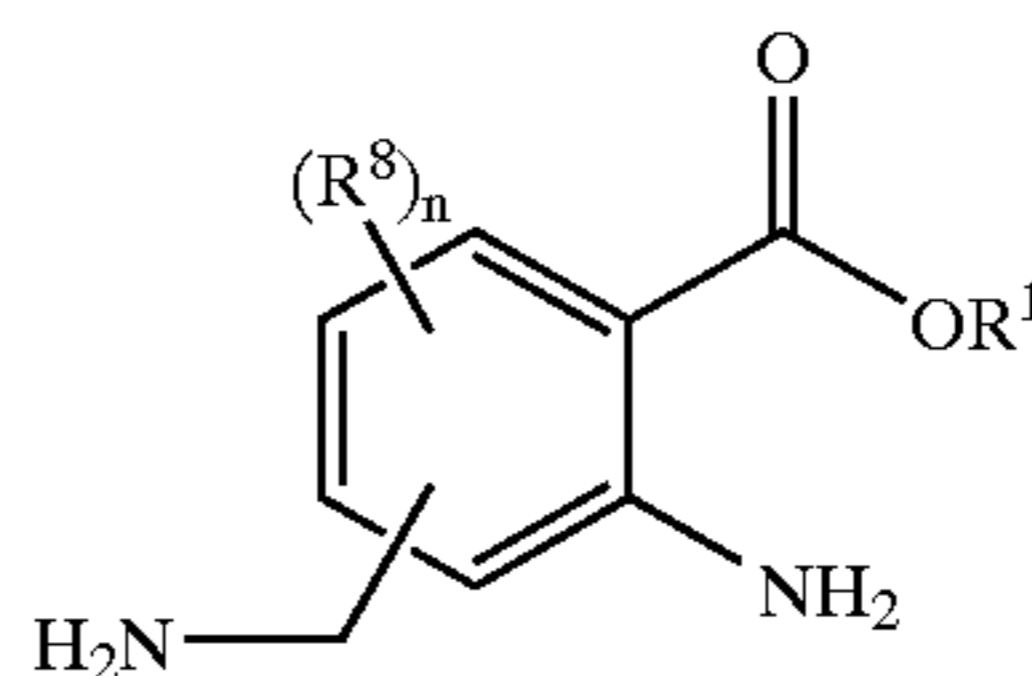
25

30

which comprises

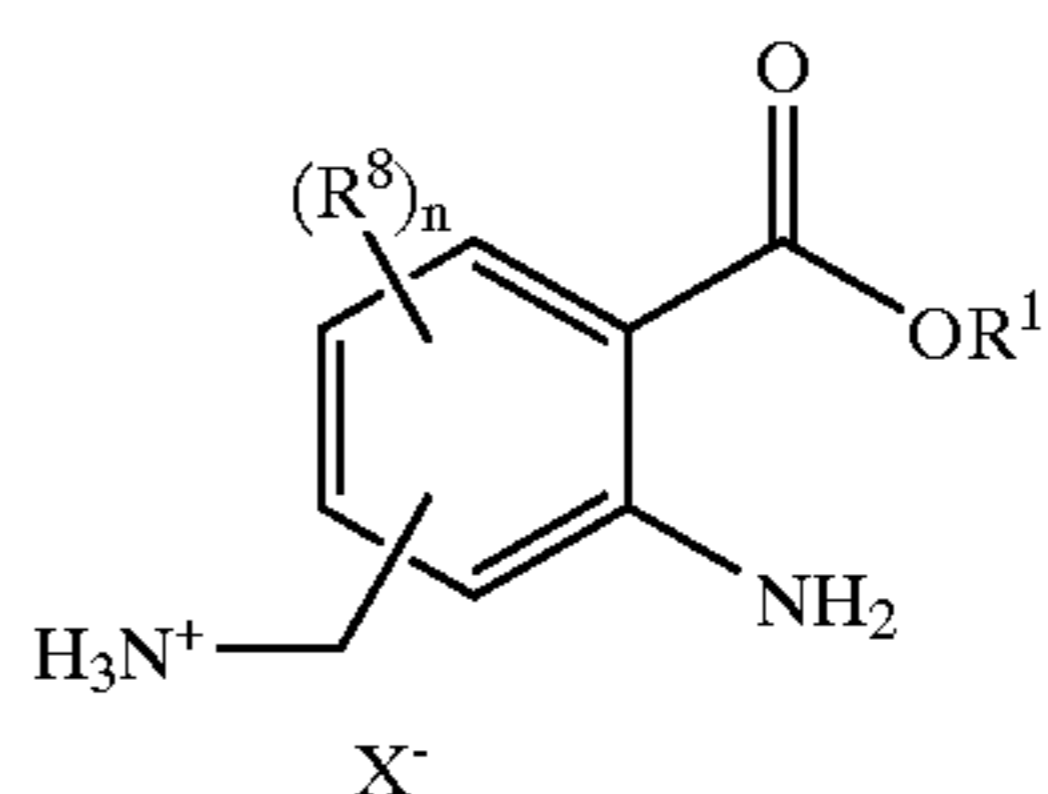
a) reacting the compound of formula (III) or (IIIa)

(III)



35

(IIIa)



45

50

with a sulfonic acid derivative to give a compound of the formula (I), wherein R¹, R², and R⁶ are hydrogen, and

and

55

b) optionally reacting a compound formed in a) with an alkylating agent or by reductive amination to give a compound of the formula (I) where R⁶ is an unsubstituted or substituted C₁-C₈-alkyl group.

* * * * *